

## Genetic Tests and Laboratory Matrix

The matrix below contains all the genetic tests which Evolent (formerly National Imaging Associates, Inc.) manages on behalf of Ambetter from Superior HealthPlan. 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.

**Refer to your [Ambetter from Superior HealthPlan clinical payment policies](#) to determine which CPT codes are allowable for each genetic test managed by Evolent.**

### Table of Contents

Table of Contents.....	1
Genetic Tests and Laboratory Matrix .....	4
Arrhythmia Panel Tests.....	4
Autism Spectrum Disorder/Intellectual Disability Panel Tests 1 .....	6
Bladder Cancer Treatment and Recurrence Algorithmic Tests .....	7
Blood-based Post Heart Transplant Gene Expression Panels for Rejection Risk.....	7
BRCA1/2 Deletion/Duplication Tests .....	8
BRCA1/2 Sequencing & Deletion/Duplication Tests.....	8

BRCA1/2 Sequencing Tests .....	10
BRCA2 Sequencing Tests.....	10
Breast Cancer Extended Endocrine Therapy Tests .....	11
Breast Cancer Prognostic Algorithmic Tests .....	12
Breast Cancer Treatment and Prognostic Algorithmic Tests .....	12
Breast DCIS Prognostic Algorithmic Tests.....	12
Cell-Free Circulating Tumor DNA Cancer Profiling Panel Tests (51 or more genes).....	12
Cell-Free Circulating Tumor DNA Cancer Profiling Panel Tests (5-50 genes) .....	13
Cell-Free Circulating Tumor DNA Lung Cancer Panel Tests .....	13
Chromosomal Microarray (SNP and CGH) for Invasive Prenatal Tests.....	14
Chromosomal Microarray (SNP and CGH) for Pregnancy Loss Tests.....	15
Chromosomal Microarray (SNP and CGH) Tests.....	17
Colorectal Cancer Prognostic Algorithmic Tests 1 .....	21
Comprehensive Arrhythmia & Cardiomyopathy (Sudden Cardiac or Unexplained Death) Panel Tests .....	21
Comprehensive Cardiomyopathy Panel Tests .....	23
Connective Tissue Disorders Panel Tests .....	24
Cutaneous Melanoma Diagnostic Algorithmic Tests .....	25
Diabetes and Obesity Panel Tests .....	25
Dilated Cardiomyopathy (DCM) Panel Tests.....	26
Donor-Derived cfDNA Testing for Heart Transplant Rejection Risk .....	27
Dystonia Panel Tests .....	27
Epilepsy and Seizure Disorder Panel Tests.....	28
Evidence-Based Lung Cancer Risk Assessment Algorithmic Tests .....	31
Evidence-Based Lung Cancer Treatment Algorithmic Tests .....	31

Evidence-Based Prostate Cancer Risk Assessment and Diagnostic Algorithmic Tests .....	31
Evidence-Based Solid Tumor Cell-free DNA (cfDNA) Panel Tests for Monitoring Minimal Residual Disease (MRD) .....	32
Exome Sequencing Tests.....	32
Expanded Carrier Panel Tests .....	38
Genome Reanalysis Tests (Interpretation Only).....	42
Genome Sequencing Tests.....	43
Hematologic Malignancy Panel Tests .....	45
Hereditary GI/Colon Cancer Panel Tests.....	48
Hereditary Polyposis Panel Tests .....	51
HLA-DQ Typing (Celiac Disease) Panel Tests.....	51
HPV-Related Solid Tumor Cell-free DNA (cfDNA) Panel Tests for Monitoring Minimal Residual Disease (MRD) .....	52
Hypertrophic Cardiomyopathy (HCM) Panel Tests.....	53
Long QT Syndrome (LQTS) Panel Tests .....	54
Lynch Syndrome / Hereditary Nonpolyposis Colorectal Cancer (HNPCC) Panel Tests .....	55
Mitochondrial Disease (including Nuclear Genes) Panel Tests.....	56
Mitochondrial DNA (mtDNA) Sequencing Panel Tests .....	58
Noonan Spectrum Disorders/RASopathies Panel Tests.....	59
Ovarian Cancer Treatment Algorithmic Tests.....	61
Pan-Cancer Hereditary Cancer Panel Tests.....	61
Pharmacogenetic Neuropsychiatric Panel Tests 1.....	64
Prenatal Cell-Free RH DNA Tests.....	65
Prostate Cancer Treatment and Prognostic Algorithmic Tests.....	65
Rapid Exome Sequencing Tests.....	66
Rapid Genome Sequencing Tests.....	67

Thyroid Cancer Diagnostic Algorithmic Tests .....	69
Tumor Specific BCR/ABL Quantitation and Breakpoint Analysis Tests.....	69
Tumor Specific Lung Cancer Panel Tests.....	71
Tumor-Type Agnostic Solid Tumor Molecular Profiling Panel Tests (51 or more genes).....	72
Tumor-Type Agnostic Solid Tumor Molecular Profiling Panel Tests (5-50 genes).....	75
Tumor-Type Agnostic Solid Tumor Molecular Profiling Panels with IHC and Cytogenetic Analyses.....	76
Uveal Melanoma Prognostic Algorithmic Tests .....	76
X-Linked Intellectual Disability Panel Tests 1.....	77

## Genetic Tests and Laboratory Matrix

### Arrhythmia Panel Tests

GTU	Test Name	Laboratory Name
<b>7VV6G</b>	Arrhythmia Comprehensive Panel	The University of Chicago Genetic Services
<b>7XHCG</b>	Arrhythmia Comprehensive Panel	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>6L93G</b>	Arrhythmia Panel	GeneDx
<b>7DMRG</b>	Arrhythmia Panel	Washington University in St. Louis Genomics and Pathology Services
<b>76V3G</b>	Atrial Fibrillation (A Fib) Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76WLG</b>	Atrial Fibrillation NGS Panel (Sequencing Only)	Fulgent Genetics
<b>5TZFG</b>	Cardiac Arrhythmia Panel	Northwest Clinical Genomics Lab
<b>5C8DG</b>	Cardiac Arrhythmia Panel	PreventionGenetics, part of Exact Sciences
<b>53VJG</b>	Comp Arrhythmia Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<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>7YKAG</b>	Comprehensive Arrhythmia Panel	Revvity
<b>2LNNG</b>	Comprehensive Arrhythmias Panel	Knight Molecular Diagnostic
<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>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>2Z2MG</b>	RhythmNext	Ambry Genetics
<b>58GWG</b>	SCA Arrhythmia Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>6DTGG</b>	Sudden Cardiac Arrest Arrhythmia Panel	GeneDx
<b>7SCDG</b>	Comprehensive Arrhythmia Gene Panel, Varies	Mayo Clinic Laboratories
<b>4ZNAG</b>	GeneSeq Cardio Familial Arrhythmia Panel	LabCorp
<b>7VA6G</b>	Arrhythmia NGS Panel	AiLife Diagnostics

GTU	Test Name	Laboratory Name
7VA7G	Arrhythmia NGS Panel Rapid	AiLife Diagnostics
7UY5G	Postmortem Arrhythmia Gene Panel, Tissue	Mayo Clinic Laboratories
7YK9G	STAT Comprehensive Arrhythmia Panel	Revvity

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

GTU	Test Name	Laboratory Name
2FHGG	Autism NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
76VUG	Autism NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
78AYG	Autism NGS Panel (Sequencing Only)	Fulgent Genetics
78FSG	Autism Spectrum Disorders (ASD) Panel	PreventionGenetics, part of Exact Sciences
7BM2G	AUTISM, INTELLECTUAL DISABILITY, and DEVELOPMENTAL DELAY GENE SEQUENCING PANEL	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
76QWG	AUTISM/AUTISM SPECTRUM DISORDER (53 gene panel)	Center for Human Genetics, Inc.
77KNG	Autism/ID Panel	GeneDx
78K3G	Autism/ID Xpanded Panel	GeneDx
77YKG	AutismNext	Ambry Genetics
78FHG	Autosomal Recessive Non-Specific Intellectual Disability Panel	The University of Chicago Genetic Services
77SAG	Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA + Fragile X Repeat Expansion and Methylation)	MNG Laboratories
77AUG	Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
76YDG	Inherited Glycosylphosphatidylinositol Biosynthesis Defects (IGDs) Panel	PreventionGenetics, part of Exact Sciences
5FW2G	Intellectual Disability Exome	The University of Chicago Genetic Services
79Q6G	Intellectual disability panel - NGS Panel (CNV included)	Centogene
775YG	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Duo (IDEA panel of patient + 1 additional family members)	PreventionGenetics, part of Exact Sciences
783YG	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Trio (IDEA panel of patient + 2 additional family members)	PreventionGenetics, part of Exact Sciences

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>77KFG</b>	Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Patient Only	PreventionGenetics, part of Exact Sciences
<b>7CFRG</b>	Invitae Neurodevelopmental Disorders (NDD) Panel	Invitae Corporation
<b>77HZG</b>	NeurodevelopmentNext	Ambry Genetics
<b>77H2G</b>	Non-Specific Intellectual Disability Panel	The University of Chicago Genetic Services
<b>76UQG</b>	Nonsyndromic Intellectual Disability (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>7BYUG</b>	PGmaxTM - Intellectual Disability, Epilepsy, and Autism (IDEA) Panel	PreventionGenetics, part of Exact Sciences
<b>7768G</b>	PGXome Custom - Intellectual Disability, Autosomal Dominant	PreventionGenetics, part of Exact Sciences
<b>77XDG</b>	PGXome Custom - Intellectual Disability, Autosomal Recessive	PreventionGenetics, part of Exact Sciences
<b>77SFG</b>	Syndromic Intellectual Disability (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>78FXG</b>	Top 99 Genetic Causes of Developmental Delay Panel	PreventionGenetics, part of Exact Sciences
<b>78F2G</b>	Comprehensive Intellectual Disability and Autism NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics

#### Bladder Cancer Treatment and Recurrence Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6W4EG</b>	Cxbladder Monitor	Pacific Edge
<b>72KHG</b>	Cxbladder Triage	Pacific Edge
<b>7BN4G</b>	Decipher Bladder Genomic Test	Veracyte
<b>7XRFG</b>	Oncuria Monitor	DiaCarta Laboratory
<b>7XREG</b>	Oncuria Predict	DiaCarta Laboratory
<b>7VLUG</b>	Cxbladder Triage Plus	Pacific Edge

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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>74CFG</b>	AlloMap Heart	Allina Health Laboratory
<b>525DG</b>	AlloMap Heart	CareDx, Inc.

## BRCA1/2 Deletion/Duplication Tests

GTU	Test Name	Laboratory Name
<b>6RPHG</b>	BRACAnalysis Rearrangement Test (BART)	Myriad Genetics
<b>79AGG</b>	BRCA1, BRCA2 panel - Deletion/Duplication Testing (MLPA/qPCR)	Centogene
<b>7QGLG</b>	BRCA1/BRCA2 DELETION/DUP BY MLPA	Clinical Pathology Laboratories
<b>4ZNUG</b>	BRCAssure: BRCA1 / 2 Deletion / Duplication Analysis	Integrated Genetics
<b>4MNDG</b>	BRCAssure: BRCA1 / 2 Deletion / Duplication Analysis	Labcorp   Oncology

## BRCA1/2 Sequencing & Deletion/Duplication Tests

GTU	Test Name	Laboratory Name
<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>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>2ELCG</b>	BRCA1 and BRCA2 Sequencing and Deletion/Duplication	Knight Molecular Diagnostic
<b>7YBUG</b>	BRCA1 and BRCA2-Associated HBOC Syndrome Panel, Sequencing and Deletion/Duplication	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>6UQAG</b>	BRCA1 and BRCA2-Associated HBOC Syndrome Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>79AHG</b>	BRCA1, BRCA2 panel Combi (with MLPA) - NGS Panel (MLPA included)	Centogene

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>79AJG</b>	BRCA1, BRCA2 panel Plus - NGS Panel (CNV included)	Centogene
<b>2YYZG</b>	BRCA1/2 seq and del/dup	Ambry Genetics
<b>6DTQG</b>	BRCA1/2 Sequencing and Del/Dup Analysis	GeneDx
<b>3D8PG</b>	BRCA1/2 Sequencing and Deletion Duplication Analysis	ACL Laboratories
<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>6LTEG</b>	BRCA1&2 Analysis	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>546VG</b>	BRCAssure: BRCA1 / 2 Comprehensive Analysis	Labcorp   Oncology
<b>58G6G</b>	BRCAssure: Comprehensive BRCA1 / 2 Analysis	Integrated Genetics
<b>6LS6G</b>	BRCAssure : BRCA1 and BRCA2 Comprehensive Analysis	LabCorp
<b>2C88G</b>	Empower - BRCA1 & BRCA2 Hereditary Cancer Test	Natera
<b>7YPXG</b>	Hereditary Breast and Ovarian Cancer Syndrome Panel	Revvity
<b>7WXHG</b>	Hereditary cancer BRCA1 BRCA2	University of Minnesota Physicians Outreach Laboratory
<b>6RMHG</b>	Integrated BRACAnalysis	Myriad Genetics
<b>44N2G</b>	Invitae BRCA1 and BRCA2 STAT Panel	Invitae Corporation
<b>6RN9G</b>	Reflex BRACAnalysis	Myriad Genetics
<b>7YPWG</b>	STAT Hereditary Breast and Ovarian Cancer Syndrome Panel	Revvity
<b>7QGJG</b>	BRCA1 AND BRCA2, COMPREHENSIVE	Clinical Pathology Laboratories
<b>7QDVG</b>	Integrated Reflex BRACAnalysis	Myriad Genetics
<b>7SLMG</b>	PrevenTest, Custom (BRCA1, BRCA2)	Advanced Molecular Diagnostics, LLC
<b>7SYQG</b>	BRAC ANALYSIS,COMPREHENSIVE	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
<b>7U27G</b>	BRCA1 and BRCA2 Sequencing and Del/Dup (NGS)	University of Michigan - Michigan Medical Genetics Laboratories
<b>442GG</b>	Invitae BRCA1 and BRCA2 Panel	Invitae Corporation
<b>7V2JG</b>	BRCA1 & BRCA2 Panel	Baylor Genetics, LLC
<b>7X27G</b>	GxVISION Hereditary Cancer Risk Assessment BRCA1/2 Genes	Otogenetics
<b>7V3GG</b>	OnkoRisk BRCA Panel (Non-NYS)	BioReference Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YDPG</b>	BRCA 1/2, COMP	ProPath
<b>57DKG</b>	Hereditary Breast, Ovarian, Pancreatic, and Prostate Cancer Syndrome BRCA1/2 Panel	PreventionGenetics, part of Exact Sciences
<b>6925G</b>	BRCA 1&2	Center for Genetic Testing at Saint Francis

### BRCA1/2 Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7QGKG</b>	BRCA1 and BRCA2 Sequencing	Clinical Pathology Laboratories
<b>6UQTG</b>	BRCA1-2 with RNAinsight	Ambry Genetics
<b>79AFG</b>	BRCA1, BRCA2 panel - NGS Panel	Centogene
<b>3SEYG</b>	BRCA1/2 Sequencing Test	Gene by Gene
<b>7U26G</b>	BRCA1 and BRCA2 Sequencing (NGS)	University of Michigan - Michigan Medical Genetics Laboratories

### BRCA2 Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6RPKG</b>	BRCA2 Analysis	Myriad Genetics
<b>2APHG</b>	BRCA2 ANALYSIS (SEQUENCING & MLPA)	Center for Human Genetics, Inc.
<b>2A8NG</b>	BRCA2 Full Gene Sequencing and Deletion/Duplication	Invitae Corporation
<b>22MXG</b>	BRCA2 Sequence Analysis	Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories
<b>7QGHG</b>	BRCA2 SEQUENCING	Clinical Pathology Laboratories
<b>4AT2G</b>	BRCA2 SEQUENCING ONLY	Center for Human Genetics, Inc.
<b>2WCCG</b>	BRCA2 Sequencing Test	Gene by Gene
<b>6U9XG</b>	BRCA2 Single Gene (Germline)	NeoGenomics Laboratories
<b>5DZFG</b>	BRCA2 Single Gene (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>4KKZG</b>	BRCA2 Single Gene (Sequencing Only)	Fulgent Genetics
<b>79DBG</b>	Breast cancer, male, susceptibility to - Carrier Testing Sanger	Centogene
<b>79DFG</b>	Breast cancer, male, susceptibility to - Sequencing + Deletion/Duplication Package	Centogene

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>79DDG</b>	Breast cancer, male, susceptibility to - Single Gene Sequencing by NGS	Centogene
<b>79DCG</b>	Breast cancer, male, susceptibility to - Single Gene Sequencing by Sanger	Centogene
<b>79DMG</b>	Breast-ovarian cancer, familial, type 2 - Carrier Testing Sanger	Centogene
<b>79DRG</b>	Breast-ovarian cancer, familial, type 2 - Sequencing + Deletion/Duplication Package	Centogene
<b>79DPG</b>	Breast-ovarian cancer, familial, type 2 - Single Gene Sequencing by NGS	Centogene
<b>79DNG</b>	Breast-ovarian cancer, familial, type 2 - Single Gene Sequencing by Sanger	Centogene
<b>79GLG</b>	Fanconi anemia type D1 - Carrier Testing Sanger	Centogene
<b>79GQG</b>	Fanconi anemia type D1 - Sequencing + Deletion/Duplication Package	Centogene
<b>79GNG</b>	Fanconi anemia type D1 - Single Gene Sequencing by NGS	Centogene
<b>79GMG</b>	Fanconi anemia type D1 - Single Gene Sequencing by Sanger	Centogene
<b>5WT9G</b>	Fanconi Anemia via the BRCA2/FANCD1 Gene	PreventionGenetics, part of Exact Sciences
<b>79J2G</b>	Glioblastoma type 3, susceptibility to, due to BRCA2 germline mutation - Carrier Testing Sanger	Centogene
<b>79J6G</b>	Glioblastoma type 3, susceptibility to, due to BRCA2 germline mutation - Sequencing + Deletion/Duplication Package	Centogene
<b>79J4G</b>	Glioblastoma type 3, susceptibility to, due to BRCA2 germline mutation - Single Gene Sequencing by NGS	Centogene
<b>79J3G</b>	Glioblastoma type 3, susceptibility to, due to BRCA2 germline mutation - Single Gene Sequencing by Sanger	Centogene

### Breast Cancer Extended Endocrine Therapy Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UU2G</b>	Breast Cancer Index	bioTheranostics
<b>759GG</b>	Breast Cancer Index (BCI)	NeoGenomics Laboratories

## Breast Cancer Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<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>6UUCG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Quest Diagnostics
<b>2KTBG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Veracyte
<b>54HTG</b>	Prosigna Breast Cancer Prognostic Gene Signature Assay	Labcorp   Oncology

## Breast Cancer Treatment and Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>2VDEG</b>	Oncotype Dx Breast Cancer Assay	Marshfield Labs
<b>7BNGG</b>	Oncotype DX Breast Recurrence Score Test	Exact Sciences Laboratories, LLC

## Breast DCIS Prognostic Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>7BNFG</b>	Oncotype DX® Breast DCIS Score	Exact Sciences Laboratories, LLC

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

GTU	Test Name	Laboratory Name
<b>7BZFG</b>	FoundationOne Liquid CDx (For Allina outpatient use only)	Allina Health Laboratory
<b>6R3DG</b>	FoundationOne Liquid CDx	Foundation Medicine
<b>7ABJG</b>	Guardant360	Guardant Health
<b>6UW6G</b>	Guardant360 CDx	Guardant Health
<b>2B64G</b>	Tempus  xF: Liquid Biopsy Panel of 105 Genes	Tempus AI, Inc.
<b>7TMMG</b>	LiquidHALLMARK	Lucence Health

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TXKG</b>	Tempus xF Liquid Biopsy Test, blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>7ACKG</b>	MSK-ACCESS	Memorial Sloan Kettering Cancer Center
<b>7VPQG</b>	Northstar Select	BillionToOne, Inc.
<b>7S8WG</b>	Tempus xF+: Liquid Biopsy Panel of 523 Genes	Tempus AI, Inc.
<b>7YXFG</b>	Caris Assure	University of Minnesota Physicians Outreach Laboratory
<b>7VVGW</b>	Caris Assure	Caris Life Sciences
<b>7Y2RG</b>	Liquid Biopsy	BostonGene
<b>7YTFG</b>	LiquidHALLMARK ctDNA and ctRNA	Mayo Clinic Laboratories
<b>7YSPG</b>	Plasma Complete	LabCorp

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

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

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

<b>4FGPG</b>	InVisionFirst -Lung Liquid Biopsy	NeoGenomics Laboratories
<b>4FGPG</b>	Resolution ctDx Lung assay (Liquid Biopsy)	NeoGenomics Laboratories
<b>7TRFG</b>	InVisionFirst -Lung Liquid Biopsy	Inivata Ltd
<b>7ACFG</b>	GeneStrat	Biodesix
<b>7ACGG</b>	GeneStrat NGS	Biodesix
<b>4764G</b>	Genestrat Test	ACL Laboratories
<b>7RVXG</b>	IQlung Treatment Guidance Testing	Biodesix

4FGPG	InVisionFirst -Lung Liquid Biopsy	NeoGenomics Laboratories
GTU	Test Name	Laboratory Name

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

GTU	Test Name	Laboratory Name
<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>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>25DWG</b>	Cytogenomic SNP Microarray, Fetal	ARUP Laboratories
<b>78KYG</b>	Cytogenomic SNP Microarray, Fetal	University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory
<b>3DQXG</b>	Cytogenomic SNP Microarray, Fetal	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>4XSNG</b>	Cytogenomic SNP Microarray, Fetal	TriCore Reference Laboratories
<b>7U5HG</b>	Cytogenomic SNP Microarray&nbsp;-&nbsp;Fetal	University of Pennsylvania School of Medicine - Molecular Pathology 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>7X43G</b>	POC Microarray with 5-Cell Chromosome Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>328BG</b>	Prenatal Cytogenomic Microarray	University Hospitals

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2LEXG</b>	Prenatal Microarray	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>7X45G</b>	Prenatal Microarray without Parental Testing	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<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>5BNSG</b>	Reveal SNP Microarray – Prenatal	Integrated Genetics
<b>2EPWG</b>	SNP Array for Prenatal Analysis (aka Microarray)	UCSF Molecular Diagnostics Laboratory
<b>7X47G</b>	SNP Microarray	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>3U6CG</b>	SNP Microarray (Direct)-Prenatal (Reveal)	LabCorp
<b>3N8VG</b>	SNP Microarray-Prenatal (Reveal)	LabCorp
<b>7X49G</b>	Prenatal Microarray with Parental Testing	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7Y5VG</b>	Prenatal chromosomal microarray analysis   CMA	Washington University in St. Louis Pathology Services
<b>7Y5TG</b>	Products of conception chromosomal microarray analysis   CMA	Washington University in St. Louis Pathology Services
<b>33X4G</b>	Chromosomal Microarray Prenatal, Amniotic fluid/Chorionic Villus Sampling (CMAP)	Marshfield Labs
<b>7V33G</b>	IriSight CNV Analysis	Variantyx, Inc.
<b>2KSUG</b>	Follow-up Prenatal CMA qPCR	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>6LRAG</b>	Chromosomal Microarray Analysis (CMA) - Products of Conception	Allina Health Laboratory
<b>7U35G</b>	Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue	University of Michigan - Michigan Medical Genetics Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6R3NG</b>	Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue	Mayo Clinic 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>2ENDG</b>	Microarray-Products of Conception (POC) Reveal FFPE	LabCorp
<b>2E65G</b>	Microarray-Products of Conception (POC) Reveal FFPE, Data Transfer	LabCorp
<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>3U6BG</b>	SNP Microarray Products of Conception (POC) / Tissue (Reveal )	LabCorp
<b>7PTBG</b>	Microarray Analysis - PREGNANCY LOSS	Nebraska Medical Center - Molecular Diagnostic Laboratory
<b>7U37G</b>	Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth	University of Michigan - Michigan Medical Genetics Laboratories
<b>6R2VG</b>	Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth	Mayo Clinic Laboratories
<b>7YEQG</b>	POC Microarray Analysis	ProPath
<b>7XJKG</b>	POC (Products of Conception) Microarray Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>2M7XG</b>	Pregnancy Loss Chromosomal SNP Microarray	Colorado Genetics Laboratory
<b>7YUVG</b>	Chromosomal Microarray, Products of Conception/Stillbirth/Varies	Hennepin County Medical Center

## 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>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>2E8LG</b>	Assure SNP Microarray Analysis (FDA cleared) – Prenatal & Postnatal	CytoGenX
<b>3GPJG</b>	Assure SNP Microarray Analysis (FDA cleared)	CytoGenX
<b>3AG7G</b>	CGH: Comparative Genomic Hybridization	New Jersey Medical School - Institute of Genomic Medicine
<b>4CURG</b>	CHROMOSOMAL MICROARRAY	Fullerton Genetics Center - Mission Health
<b>7YYMG</b>	CHROMOSOMAL MICROARRAY	DDC Clinic Laboratory
<b>3F5GG</b>	CHROMOSOMAL MICROARRAY	Detroit Medical Center University Laboratories - Molecular Genetics Diagnostic Laboratory
<b>38VKG</b>	Chromosomal Microarray Analysis	UCLA Diagnostic Molecular Pathology Laboratory
<b>687XG</b>	Chromosomal Microarray Analysis	Center for Genetic Testing at Saint Francis
<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>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
<b>333GG</b>	Chromosomal Microarray with reflex karyotype	Washington University in St. Louis Pathology Services
<b>6R2TG</b>	Chromosomal Microarray, Congenital, Blood	Mayo Clinic Laboratories
<b>7YUWG</b>	Chromosomal Microarray, Congenital, Blood	Hennepin County Medical Center
<b>2Z55G</b>	Chromosomal Microarray, POC FFPE, ClariSure Oligo-SNP	Quest Diagnostics
<b>8EJVG</b>	Chromosomal Microarray, POC, ClariSure Oligo-SNP	Medfusion

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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	Johns Hopkins All Children's Hospital
<b>2DYKG</b>	CHROMOSOME MICROARRAY, POSTNATAL	Empire City Laboratories, Inc.
<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>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
<b>2ETHG</b>	Constitutional SNP Array Karyotyping, Constitutional Chromosome Microarray Analysis (CMA)	Beaumont Laboratories - Molecular Pathology Lab
<b>2M4WG</b>	Cytogenomic Microarray Analysis	University of Washington Medicine - Pathology - Cytogenetics and Genomics Laboratory
<b>49JCG</b>	Cytogenomic Microarray Analysis of Postnatal Blood	Akron Children's Hospital
<b>49W6G</b>	Cytogenomic SNP array (postnatal)	The University of Chicago Genetic Services
<b>2MTFG</b>	Cytogenomic SNP Microarray	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<b>7XL3G</b>	Cytogenomic SNP Microarray	University of Illinois at Chicago - Biochemical Genetics Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2X3MG</b>	Cytogenomic SNP Microarray	University of California Davis Health System - UC DMC - Molecular and Cytogenetic Laboratory
<b>8EBHG</b>	Cytogenomic SNP Microarray	Medfusion
<b>2E3BG</b>	Cytogenomic SNP Microarray	ARUP Laboratories
<b>7U9JG</b>	Cytogenomic SNP Microarray	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>2542G</b>	Cytogenomic SNP Microarray Buccal Swab	ARUP Laboratories
<b>3MA8G</b>	Genomic Microarray, Blood	ACL Laboratories
<b>2ENHG</b>	Genomic SNP Microarray, Products of Conception	University of California Davis Health System - UC DMC - Molecular and Cytogenetic Laboratory
<b>7UHKG</b>	Genomic SNP Microarray, Products of Conception	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>2ETGG</b>	Illumina Global Screening Array Sequencing Test	Gene by Gene
<b>2EQNG</b>	Illumina Microarray Analysis	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
<b>2EMZG</b>	Illumina Microarray Analysis- Targeted Family	University of Wisconsin - Madison WSLH - UW Cytogenetic Services
<b>2EMYG</b>	Invitae Chromosomal Microarray Analysis (CMA) with follow-up FISH when required	Invitae Corporation
<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>3LQLG</b>	Microarray Analysis - CytoScan SNP	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
<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>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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2EKFG</b>	Reflex to SNP Array	UCSF Molecular Diagnostics Laboratory
<b>2DYMG</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
<b>2Z5JG</b>	SNP Microarray Pediatric	Integrated Regional Laboratories
<b>2EKEG</b>	SNP microarray Processing, Extraction and Storage	UCSF Molecular Diagnostics Laboratory
<b>49AAG</b>	SNP Microarray when Routine Chrom or High Res and Fragile X have ALREADY been billed	Center for Human Genetics, Inc.
<b>52MEG</b>	SNP Oligonucleotide Microarray Analysis (SOMA)	Columbia University - Personalized Genomic Medicine
<b>6NNKG</b>	Whole-Genome Chromosomal Microarray (CMA-ISCA) via the aCGH and SNP Test	PreventionGenetics, part of Exact Sciences
<b>7QHSG</b>	CHROMOSOME SNP MICROARRAY	University of Texas Medical Branch - UTMB - Porphyria Laboratory
<b>7QHGG</b>	LV CYTOGENOMIC SNP MICROARRAY	Clinical Pathology Laboratories
<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>4DNFG</b>	Chromosomal Microarray Analysis for Constitutional Abnormalities	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>7R9LG</b>	Chromosomal Microarray, Hematologic Malignancy, ClariSure Oligo-SNP (90961)	Rady Children's Institute for Genomic Medicine
<b>7WJHG</b>	Constitutional Chromosomal Microarray (Copy Number)	University of Minnesota Physicians Outreach Laboratory
<b>7WPQG</b>	Constitutional or Products of Conception (POC) Chromosomal Microarray (Copy Number/SNP)	University of Minnesota Physicians Outreach Laboratory
<b>2EQQG</b>	SNP Microarray-Pediatric (Reveal)	LabCorp
<b>6XNCG</b>	Chromosomal Microarray (MicroarrayDx)	GeneDx
<b>7S97G</b>	Chromosome Microarray, Congenital	Sanford USD Medical Center - Sanford Clinic USD Genetics Laboratory

GTU	Test Name	Laboratory Name
7WBMG	Constitutional Limited Chromosomal Microarray (Copy number only) (Charged)	University of Minnesota Physicians Outreach Laboratory
7TG6G	Invitae Chromosomal Microarray Analysis (CMA)	Invitae Corporation
7UHHG	Genomic Microarray Analysis - Parental study, Genome Wide Array	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
7W9QG	Cytogenomic Microarray SNP Fetal	University of Minnesota Physicians Outreach Laboratory
7XJLG	Autopsy Microarray Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
7X4CG	CNS Tumor Classification by Methylation Array	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
2DYJG	Postnatal Chromosomal SNP Microarray	Colorado Genetics Laboratory
7WXWG	SNP Array for Cancer Analysis	UCSF Molecular Diagnostics Laboratory
7YHLG	Chromosomal Microarray, Congenital, Blood (CMACB)	Marshfield Labs
7YTUG	Genomic Array, FHCC	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory

#### 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
2LRSG	Combined Cardiac Panel	GeneDx
23DQG	Cardiomyopathy and Arrhythmia Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
2DTLG	Cardiomyopathy and Arrhythmia Sequencing Panel	UCSD - Molecular Diagnostics & Cytogenetics Laboratory
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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>43H3G</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy	Invitae Corporation
<b>43FSG</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes	Invitae Corporation
<b>7UY6G</b>	Postmortem Cardiomyopathy and Arrhythmia Gene Panel, Tissue	Mayo Clinic Laboratories
<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
<b>7VV4G</b>	Arrhythmia and Cardiomyopathy Comprehensive Panel	The University of Chicago Genetic Services
<b>7TP5G</b>	CardioNext: Arrhythmia	Ambry Genetics
<b>77BXG</b>	CardioNext: Cardiomyopathy	Ambry Genetics
<b>7DR2G</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel - UNLOCKCARDIO	Invitae Corporation
<b>7QKVG</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy - UNLOCKCARDIO	Invitae Corporation
<b>7QL3G</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes - UNLOCKCARDIO	Invitae Corporation
<b>7YWEG</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel - ROC01	Invitae Corporation
<b>7YWAG</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy - ROC01	Invitae Corporation
<b>7YWDG</b>	Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes - ROC01	Invitae Corporation
<b>78FVG</b>	Comprehensive Arrhythmia and Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>7YNRG</b>	WholeCardiologyTM Panel	Revvity

## Comprehensive Cardiomyopathy Panel Tests

GTU	Test Name	Laboratory Name
<b>7DMVG</b>	Cardiomyopathy Panel	Washington University in St. Louis Genomics and Pathology Services
<b>4MUVG</b>	Cardiomyopathy Panel	Northwest Clinical Genomics Lab
<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>3XZHG</b>	Comprehensive Cardiomyopathy NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<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>7YKEG</b>	Comprehensive Cardiomyopathy Panel	Revvity
<b>3GBZG</b>	GeneSeq : Cardio-Familial Cardiomyopathy Profile	Integrated Genetics
<b>5FWFG</b>	Invitae Cardiomyopathy Comprehensive Panel	Invitae Corporation
<b>24WTG</b>	Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy	Invitae Corporation
<b>7BLQG</b>	Pan Cardiomyopathy Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>57C9G</b>	Pan Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>7YKDG</b>	STAT Comprehensive Cardiomyopathy Panel	Revvity
<b>7SCCG</b>	Comprehensive Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>56ZPG</b>	GeneSeq Cardio Familial Cardiomyopathy Panel	LabCorp
<b>7UHGG</b>	Pan Cardiomyopathy Panel (62 Genes)	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>7VAEG</b>	Cardiomyopathy NGS Panel	AiLife Diagnostics

GTU	Test Name	Laboratory Name
7VAFG	Cardiomyopathy NGS Panel Rapid	AiLife Diagnostics
7UY7G	Postmortem Cardiomyopathy Gene Panel, Tissue	Mayo Clinic Laboratories
3YNXG	Invitae Cardiomyopathy Comprehensive Panel - LYSO	Invitae Corporation
3YRRG	Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy - LYSO	Invitae Corporation
7YH2G	Invitae Cardiomyopathy Comprehensive Panel - ROC01	Invitae Corporation
7YH3G	Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy - ROC01	Invitae Corporation

### Connective Tissue Disorders Panel Tests

GTU	Test Name	Laboratory Name
5LFXG	Brittle Cornea Syndrome Panel	PreventionGenetics, part of Exact Sciences
3QTPG	C1S and C1R gDNA Testing	Collagen Diagnostic Laboratory
78ADG	CONNECT2: CONNECTIVE TISSUE DISORDERS DNA SEQUENCING CHIP	Center for Human Genetics, Inc.
79PZG	Connective tissue and related disorders panel - NGS Panel (CNV included)	Centogene
7VANG	Connective Tissue Disorder NGS Panel	AiLife Diagnostics
7VAPG	Connective Tissue Disorder NGS Panel Rapid	AiLife Diagnostics
789VG	Connective Tissue Disorder Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
7725G	Connective Tissue Disorders NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
77MRG	Connective Tissue Disorders Panel	PreventionGenetics, part of Exact Sciences
7VXMG	Connective Tissue Disorders Panel	Baylor Genetics, LLC
4VXTG	Connective Tissue NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
77FCG	Connective Tissue NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
77DJG	Connective Tissue NGS Panel (Sequencing Only)	Fulgent Genetics
76QEG	Connective Tissue Panel	Knight Molecular Diagnostic
77MDG	Heritable Disorders of Connective Tissue Panel	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7875G</b>	Invitae Connective Tissue Disorders Panel	Invitae Corporation
<b>77YXG</b>	Skin and Connective Tissue Disorders Panel	PreventionGenetics, part of Exact Sciences

### Cutaneous Melanoma Diagnostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7BR6G</b>	DecisionDx DiffDx-Melanoma	Castle Biosciences Inc
<b>7BR5G</b>	myPath Melanoma	Castle Biosciences Inc

### Diabetes and Obesity Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>78LNG</b>	Comprehensive Monogenic Obesity Panel	PreventionGenetics, part of Exact Sciences
<b>79Q2G</b>	Diabetes and obesity panel - NGS Panel (CNV included)	Centogene
<b>679FG</b>	Diabetes-Obesity NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>77TVG</b>	Diabetes-Obesity NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>78D4G</b>	Diabetes-Obesity NGS Panel (Sequencing Only)	Fulgent Genetics
<b>7VASG</b>	Diabetes/MODY and Obesity NGS Panel	AiLife Diagnostics
<b>7VATG</b>	Diabetes/MODY and Obesity NGS Panel Rapid	AiLife Diagnostics
<b>47Q5G</b>	Early-Onset Obesity Evaluation	Athena Diagnostics Inc
<b>782ZG</b>	Invitae Monogenic Diabetes Panel	Invitae Corporation
<b>76TCG</b>	Invitae Monogenic Obesity Panel	Invitae Corporation
<b>77YYG</b>	Monogenic Diabetes Panel	PreventionGenetics, part of Exact Sciences
<b>77H8G</b>	Monogenic Obesity Panel	The University of Chicago Genetic Services
<b>78C6G</b>	No-charge Sponsored Testing Program for Rare Genetic Diseases of Obesity	PreventionGenetics, part of Exact Sciences
<b>78LRG</b>	Non-Syndromic Monogenic Obesity Panel	PreventionGenetics, part of Exact Sciences
<b>77H9G</b>	Non-Syndromic Monogenic Obesity Panel	The University of Chicago Genetic Services
<b>38G6G</b>	Suggested Custom Slice - Obesity	GeneDx

## Dilated Cardiomyopathy (DCM) Panel Tests

GTU	Test Name	Laboratory Name
<b>2YW9G</b>	DCMNext	Ambry 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>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>7XHEG</b>	Dilated Cardiomyopathy Panel	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>3CBUG</b>	Dilated Cardiomyopathy Panel	Knight Molecular Diagnostic
<b>5CZ6G</b>	Dilated Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<b>7YKCG</b>	Dilated Cardiomyopathy Panel	Revuity
<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>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>7YKBG</b>	STAT Dilated Cardiomyopathy Panel	Revuity
<b>5NAGG</b>	Dilated & Arrhythmogenic Cardiomyopathy NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<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

## Donor-Derived cfDNA Testing for Heart Transplant Rejection Risk

GTU	Test Name	Laboratory Name
<b>7R84G</b>	AlloSureHeart	CareDx, Inc.
<b>86PMG</b>	Prospera Heart	Natera
<b>7UVUG</b>	Viracor TRAC® Heart dd-cfDNA	Eurofins Viracor, LLC

## 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>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>775BG</b>	Dystonia Panel	Knight Molecular Diagnostic
<b>775XG</b>	Dystonia Panel	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>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

### 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>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>78L9G</b>	Clinical Epilepsy NGS Panel	MNG Laboratories
<b>87ASG</b>	Clinical Epilepsy NGS Panel	LabCorp
<b>77ASG</b>	Comprehensive Epilepsy (NGS Panel and Copy Number Analysis + mtDNA)	MNG Laboratories
<b>73X4G</b>	Comprehensive Epilepsy NGS Panel	LabCorp
<b>78KDG</b>	Comprehensive Epilepsy Panel	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YKZG</b>	Comprehensive Epilepsy Panel	Revvity
<b>7BNUG</b>	Comprehensive Epilepsy With or Without Encephalopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>78FGG</b>	Early Infantile Epileptic Encephalopathy (EIEE) Panel	The University of Chicago Genetic Services
<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>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>77GZG</b>	Epilepsy Exome	The University of Chicago Genetic Services
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>78FEG</b>	Epilepsy Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7XHKG</b>	Epilepsy Panel	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7V2RG</b>	Epilepsy Panel	Baylor Genetics, LLC
<b>79Q4G</b>	Epilepsy panel - NGS Panel (CNV included)	Centogene
<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>775EG</b>	EpiXpanded Panel	GeneDx
<b>77P9G</b>	Infantile Epilepsy Panel	Knight Molecular Diagnostic
<b>3Q7DG</b>	Infantile Spasms NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<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>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>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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>78EUG</b>	PGXome Custom - Early Infantile Epileptic Encephalopathy, Recessive	PreventionGenetics, part of Exact Sciences
<b>7YKYG</b>	STAT Comprehensive Epilepsy Panel	Revvity
<b>7V2QG</b>	STAT Epilepsy Panel	Baylor Genetics, LLC
<b>77Y5G</b>	Epilepsy/Seizure NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>6YMKG</b>	Genomic Unity Epilepsy Analysis	Variantyx, Inc.
<b>7UUBG</b>	Comprehensive Epilepsy Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>78HSG</b>	PGmaxTM - Comprehensive Epilepsy and Seizure Panel	PreventionGenetics, part of Exact Sciences
<b>7WZ6G</b>	Invitae Epilepsy Panel - UNLOCKBTS	Invitae Corporation
<b>7WZ7G</b>	Invitae Epilepsy Panel-Add-on Preliminary-evidence Genes for Epilepsy - UNLOCKBTS	Invitae Corporation

#### Evidence-Based Lung Cancer Risk Assessment Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6USVG</b>	Nodify XL2	Biodesix

#### Evidence-Based Lung Cancer Treatment Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YCWG</b>	RiskReveal	Razor Genomics
<b>6UT9G</b>	VeriStrat	Biodesix

#### Evidence-Based Prostate Cancer Risk Assessment and Diagnostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UTPG</b>	4K Prostate Score (Serum)	BioReference Laboratories
<b>2MZ5G</b>	ConfirmMDx for Prostate Cancer	MDx Health
<b>747YG</b>	ExoDx Prostate Test	ExosomeDx
<b>7UN9G</b>	IsoPSA	Cleveland Diagnostics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7SNJG</b>	MyProstateScore 2.0	LynxDx
<b>5QLDG</b>	Prostate Cancer Gene 3	Integrated Regional Laboratories
<b>6UTAG</b>	Prostate-Specific Kallikrein, 4Kscore	ARUP Laboratories
<b>2LQFG</b>	SelectMDx for Prostate Cancer	MDx Health

### Evidence-Based Solid Tumor Cell-free DNA (cfDNA) Panel Tests for Monitoring Minimal Residual Disease (MRD)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>86PHG</b>	Guardant360 Response	Guardant Health
<b>86PKG</b>	GuardantReveal	Guardant Health
<b>6UNFG</b>	Signatera - Residual Disease Test (MRD)	Natera

### Exome Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>79AQG</b>	CentoDx - NGS Panel	Centogene
<b>46WXG</b>	CHOP Medical Exome + MitoGenome Combined Test	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<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>7XYWG</b>	Clinical Exome (Sequencing) Trio	DNA Diagnostic Laboratory - Johns Hopkins Hospital
<b>7BMXG</b>	Clinical Exome (Sequencing) Trio	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>24C9G</b>	Clinical Exome Sequencing - Duo (Proband and 1 family member)	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>86SUG</b>	Duo Whole Exome Sequencing	Baylor Genetics, LLC

GTU	Test Name	Laboratory Name
<b>3V27G</b>	Exome Select	The University of Chicago Genetic Services
<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>2WWZG</b>	Exome Sequencing Re-analysis	Northwest Clinical Genomics Lab
<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>28FHG</b>	ExomeNext-Duo plus mtDNA	Ambry Genetics
<b>2YZMG</b>	ExomeNext-Duo	Ambry Genetics
<b>2YZQG</b>	ExomeNext-Proband plus mtDNA	Ambry Genetics
<b>2YZRG</b>	ExomeNext-Proband	Ambry Genetics
<b>5U8XG</b>	ExomeNext-Trio plus mtDNA	Ambry Genetics
<b>2YZVG</b>	ExomeNext-Trio	Ambry Genetics
<b>33Z3G</b>	ExomeSeq - Whole Exome Sequencing	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7A7AG</b>	Family Member Comparator Specimen for Exome Sequencing, Varies	Mayo Clinic Laboratories
<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>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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>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>5E3RG</b>	PGxome Diagnostic Exome Test Patient Only	PreventionGenetics, part of Exact Sciences
<b>6LUFG</b>	PGxome Prenatal Exome Test - Duo	PreventionGenetics, part of Exact Sciences
<b>5NGYG</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>6QZHG</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>2ZBFG</b>	Proband Whole Exome Sequencing	Baylor Genetics, LLC
<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>2G7GG</b>	Sequential Trio Whole Exome Sequencing	Baylor Genetics, LLC
<b>6QQMG</b>	Trio Whole Exome Sequencing	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>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>6QZFG</b>	Whole Exome Sequencing	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>2Z7LG</b>	Whole Exome Sequencing	Gene by Gene
<b>7RSXG</b>	Whole Exome Sequencing	Rady Children's Institute for Genomic Medicine

Centene TX Ambetter Superior Genetic Testing and Laboratory Matrix 09/1/2025.docx  
 9/1/2025

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7R8CG</b>	Whole Exome Sequencing	Washington University in St. Louis Genomics and Pathology Services
<b>7WZRG</b>	Whole Exome Sequencing	Otogenetics
<b>5JQPG</b>	Whole Exome Sequencing - DUO (Proband)	LabCorp
<b>5JR8G</b>	Whole Exome Sequencing - Proband Only	LabCorp
<b>7YMAG</b>	Whole Exome Sequencing - Proband Only	Revvity
<b>5XKLG</b>	Whole Exome Sequencing - TRIO (Proband)	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>7YLTG</b>	Whole Exome Sequencing DATA Only (per sample)	Revvity
<b>7A7EG</b>	Whole Exome Sequencing for Hereditary Disorders , Varies	Mayo Clinic Laboratories
<b>7YLZG</b>	Whole Exome Sequencing QUAD	Revvity
<b>7YM7G</b>	Whole Exome Sequencing TRIO	Revvity
<b>7YM4G</b>	Whole Exome Sequencing, DUO	Revvity
<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>46N6G</b>	XomeDx Trio	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>6L9SG</b>	XomeDx - Proband	GeneDx
<b>6RVYG</b>	XomeDx - Trio	GeneDx
<b>4JJWG</b>	XomeDx Fetal - Duo	GeneDx
<b>7DQWG</b>	XomeDx Plus - Duo	GeneDx
<b>77MHG</b>	XomeDx Plus - Proband	GeneDx

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>77M2G</b>	XomeDx Plus - Trio	GeneDx
<b>2W48G</b>	XomeDx Prenatal - Comprehensive	GeneDx
<b>4HM8G</b>	XomeDxFetal- Proband	GeneDx
<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>7PQEG</b>	Whole Exome Sequencing – DUO (Proband), Products of Conception (POC)	LabCorp
<b>7PQFG</b>	Whole Exome Sequencing – Proband Only, Products of Conception (POC)	LabCorp
<b>7PQDG</b>	Whole Exome Sequencing – TRIO (Proband), Products of Conception (POC)	LabCorp
<b>7ABTG</b>	Genomic Unity Exome Plus Analysis	Variantyx, Inc.
<b>6L9QG</b>	XomeDx - Duo	GeneDx
<b>46YFG</b>	CHOP Medical Exome	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7TLGG</b>	Expanded Exome Upgrade per Sample	Praxis Genomics, LLC
<b>7TLMG</b>	Expanded Exome, 4 Sample	Praxis Genomics, LLC
<b>7TLKG</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>6ZSHG</b>	Genomic Unity Exome Analysis	Variantyx, Inc.
<b>2BFEG</b>	UCSF Genomics Blood Draw	UCSF Molecular Diagnostics Laboratory
<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>7TYRG</b>	Critical Trio Whole Exome Sequencing	University of Michigan - Michigan Medical Genetics Laboratories
<b>7XA6G</b>	Critical Trio Whole Exome Sequencing	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory

<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>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>7TCVG</b>	XomeDX Plus-Trio, buccal kit	University of Michigan - Michigan Medical Genetics Laboratories
<b>7YMNG</b>	Add Familial Report to Previous Whole Exome Sequencing TRIO Test	Revvity
<b>7UNTG</b>	Exome Sequencing	ARUP Laboratories
<b>7VSBG</b>	Exome Sequencing	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<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>776NG</b>	Trio Whole Exome Sequencing + Comprehensive mtDNA Analysis	Baylor Genetics, LLC
<b>2LNHG</b>	WES - Additional Affected Sibling	Baylor Genetics, LLC
<b>7VBYG</b>	Duo Exome	AiLife Diagnostics
<b>7VBUG</b>	Proband Exome	AiLife Diagnostics
<b>7VBWG</b>	Trio Exome	AiLife Diagnostics
<b>7TVKG</b>	XomeDx Express (GeneDx), Blood	University of Michigan - Michigan Medical Genetics Laboratories
<b>7VXCG</b>	Whole Exome Sequencing Reflex	Baylor Genetics, LLC
<b>7WADG</b>	Hereditary Exome Family Member	University of Minnesota Physicians Outreach Laboratory
<b>7XYZG</b>	Clinical Exome Sequencing (Duo)	DNA Diagnostic Laboratory - Johns Hopkins Hospital
<b>7BMKG</b>	Clinical Exome Sequencing (Duo)	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7BMVG</b>	Clinical Exome Sequencing (Proband Only)	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7XYYG</b>	Clinical Exome Sequencing (Proband Only)	DNA Diagnostic Laboratory - Johns Hopkins Hospital
<b>7XYXG</b>	Clinical Exome Sequencing (Quad)	DNA Diagnostic Laboratory - Johns Hopkins Hospital
<b>7BMWYG</b>	Clinical Exome Sequencing (Quad)	Johns Hopkins Medical Institutions - Pathology Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7WZHG</b>	Whole Exome with RISE Analysis (RNA-Seq) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7XAXG</b>	Rapid Whole Exome Sequencing Patient Only	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7X4AG</b>	Somatic Disease/Germline Comparator Exome	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7WZGG</b>	Whole Exome with RISE Analysis (RNA-Seq) (Sequencing Only)	Fulgent Genetics
<b>7XW6G</b>	ExomeReveal	Ambry Genetics
<b>7XW3G</b>	GeneDx Epilepsy Partnership – XomeDx - Duo	GeneDx
<b>7XW4G</b>	GeneDx Epilepsy Partnership – XomeDx - Proband	GeneDx
<b>7XW5G</b>	GeneDx Epilepsy Partnership – XomeDx - Trio	GeneDx
<b>7DQDG</b>	Whole Exome and Mitochondrial Genome Sequencing, Varies	Mayo Clinic Laboratories
<b>7WKBG</b>	XomeDx to GeneDx	University of Minnesota Physicians Outreach Laboratory
<b>7Y9GG</b>	Columbia Diagnostic Exome (CDEX)	Columbia University - Personalized Genomic Medicine
<b>7YA5G</b>	Exome Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>7YA4G</b>	Panel Reflex to Exome Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>7YANG</b>	Xenome ID	Advanced Molecular Diagnostics, LLC
<b>7YAMG</b>	Xenome Plus	Advanced Molecular Diagnostics, LLC
<b>7YLUG</b>	Reflex to Whole Exome Sequencing TRIO (from WES Proband)	Revvity
<b>7YLWG</b>	Whole Exome Sequencing, Quint	Revvity
<b>7YSDG</b>	Panel to Whole Exome Sequencing Reflex Test, Varies	Mayo Clinic Laboratories
<b>5TQ3G</b>	Invitae Exome, Trio	Invitae Corporation
<b>7Z2WG</b>	MVL Whole Exome Sequencing	Molecular Vision Laboratory

### Expanded Carrier Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>7V27G</b>	GeneAware - Complete Panel - Female	Baylor Genetics, LLC
<b>7V22G</b>	GeneAware - Complete 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>77ENG</b>	Inheritest Carrier Screen - Society Guided Panel (14 Genes)	Integrated Genetics
<b>77LMG</b>	Inheritest 500 PLUS Panel	Integrated Genetics
<b>77BTG</b>	Inheritest 500 PLUS Panel	LabCorp
<b>7838G</b>	Inheritest 500 PLUS with Repro Partners Report	Integrated Genetics
<b>7ZPEG</b>	MYRIAD FORESIGHT CARRIER SCREENING	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>3EHWG</b>	NewbornGenID	Advanced Molecular Diagnostics, LLC
<b>2D58G</b>	NewbornGenID include Fragile X (Female Patients Only)	Advanced Molecular Diagnostics, LLC
<b>6UVXG</b>	QHerit; Expanded Carrier Screen	Quest Diagnostics
<b>7VXSG</b>	QHerit; Expanded Carrier Screen	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7DHGG</b>	QHerit; Extended, Female	Quest Diagnostics
<b>7DHHG</b>	QHerit; Extended, Male	Quest Diagnostics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7DHJG</b>	QHerit; Plus, Female	Quest Diagnostics
<b>7DHKG</b>	QHerit; Plus, Male	Quest Diagnostics
<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>6RN4G</b>	Foresight Fundamental Plus panel	Myriad Genetics
<b>78HYG</b>	Foresight Universal Panel Carrier Screen	Myriad Genetics
<b>784RG</b>	MYRIAD WOMENS HEALTH FAMILY PREP SCREEN 2	University of California Davis Health System - UC DMC - Molecular and Cytogenetic Laboratory
<b>7SRSG</b>	M Beacon Focus B, 14 Gene Panel	LabCorp
<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>7SUSG</b>	Inheritest 100 PLUS Panel	LabCorp
<b>7SUTG</b>	Inheritest 300 PLUS Panel	LabCorp
<b>7SURG</b>	Inheritest High Frequency Panel	LabCorp
<b>7SUAG</b>	QHerit; 381 Diseases, Male	Quest Diagnostics
<b>7SU9G</b>	QHerit; 421 Diseases, Female	Quest Diagnostics
<b>7V3RG</b>	Comprehensive Carrier Screening Panel for Genetic Conditions	Genesys Diagnostics Inc
<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>7VXPG</b>	GeneAware Expanded Plus Panel (Female)	Baylor Genetics, LLC
<b>7VXQG</b>	GeneAware Expanded Plus Panel (Male)	Baylor Genetics, LLC
<b>7VXWG</b>	QHerit; 400 Diseases, Male	Quest Diagnostics
<b>7VXXG</b>	QHerit; 445 Diseases, Female	Quest Diagnostics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7WZBG</b>	Beacon Preconception 787-Expanded Carrier Screening Panel (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7WZCG</b>	Beacon Preconception 787-Expanded Carrier Screening Panel (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7X2CG</b>	GxVISION Carrier Screening Test ACOG/ACMG with CF	Otogenetics
<b>7X29G</b>	GxVISION Carrier Screening Test Pan-Ethnic Carrier Screening	Otogenetics
<b>7WZ9G</b>	Beacon Preconception Expanded Carrier Screening (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7WZAG</b>	Beacon Preconception Expanded Carrier Screening (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7XWVG</b>	QHerit; 4 Diseases, Male	Quest Diagnostics
<b>7XWUG</b>	QHerit; 6 Diseases, Female	Quest Diagnostics
<b>7YQUG</b>	QHerit; 112 Diseases, Female	Quest Diagnostics
<b>7YQTG</b>	QHerit; 179 Diseases, Male	Quest Diagnostics
<b>7YQSG</b>	QHerit; 198 Diseases, Female	Quest Diagnostics
<b>7YQRG</b>	QHerit; 559 Diseases, Male	Quest Diagnostics
<b>7YQQG</b>	QHerit; 611 Diseases, Female	Quest Diagnostics
<b>7YQPG</b>	QHerit; 96 Diseases, Male	Quest Diagnostics
<b>6LMNG</b>	Beacon Core Guidelines Carrier Screening Panel (With X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>5DRBG</b>	Beacon Core Guidelines Carrier Screening Panel (Without X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>78HAG</b>	Beacon Expanded Carrier Screening Panel (With X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77ZJG</b>	Beacon Expanded Carrier Screening Panel (Without X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>77ZHG</b>	Beacon Expanded Carrier Screening Plus Panel (With X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7743G</b>	Beacon Expanded Carrier Screening Plus Panel (Without X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>86GSG</b>	Beacon High Frequency Carrier Screening (With X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>86GTG</b>	Beacon High Frequency Carrier Screening (Without X-linked Disorders) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7WZDG</b>	Beacon Preconception High Frequency Carrier Screening (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7WZEG</b>	Beacon Preconception High Frequency Carrier Screening (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication)	Fulgent Genetics

### Genome Reanalysis Tests (Interpretation Only)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YNLG</b>	Analysis and Interpretation of Whole Genome Sequencing External Data	Revvity
<b>7TL3G</b>	External Data Analysis, 4 Sample	Praxis Genomics, LLC
<b>7TKZG</b>	External Data Analysis, Duo	Praxis Genomics, LLC
<b>7TKYG</b>	External Data Analysis, Proband	Praxis Genomics, LLC
<b>7TL2G</b>	External Data Analysis, Trio	Praxis Genomics, LLC
<b>2WWMG</b>	GenomeSeqDx Reanalysis	GeneDx
<b>7YMUG</b>	Reanalysis and Interpretation of Previous RVTY Whole Genome DUO Sequencing Test	Revuity
<b>7YMGV</b>	Reanalysis and Interpretation of Previous RVTY Whole Genome Proband Sequencing Test	Revuity
<b>7YMSG</b>	Reanalysis and Interpretation of Previous RVTY Whole Genome QUAD Sequencing Test	Revuity
<b>7YMTG</b>	Reanalysis and Interpretation of Previous RVTY Whole Genome TRIO Sequencing Test	Revuity
<b>7XCAG</b>	Reanalysis Genome Sequencing-Parent Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7V2YG</b>	WGS Reanalysis	Baylor Genetics, LLC
<b>7T4HG</b>	Whole Genome Reanalysis	ARUP Laboratories
<b>7SAUG</b>	Whole Genome Sequencing Reanalysis, Varies	Mayo Clinic Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YNMG</b>	Analysis and Interpretation for Whole Exome Sequencing External Data	Revvity
<b>7YMPG</b>	First Reanalysis on Previous RVTY Whole Genome Sequencing Test	Revvity
<b>7YN2G</b>	Reanalysis and Interpretation of Previous RVTY Whole Exome Proband Sequencing Test	Revvity
<b>7YMRG</b>	Reanalysis and Interpretation of Previous RVTY Whole Genome QUINT Sequencing Test	Revvity
<b>2LUBG</b>	Genome Sequencing Subsequent Reanalysis (charged)	GeneDx
<b>7XCBG</b>	Reanalysis Genome Sequencing-Patient Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory

### Genome Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YMMG</b>	Add Familial Report to Previous Whole Genome Sequencing TRIO Test	Revvity
<b>7V9YG</b>	Duo WGS	AiLife Diagnostics
<b>86SWG</b>	Duo Whole Genome Sequencing	Baylor Genetics, LLC
<b>7SC9G</b>	Family Member Comparator Specimen for Genome Sequencing, Varies	Mayo Clinic Laboratories
<b>7TVEG</b>	Genome Express	University of Michigan - Michigan Medical Genetics Laboratories
<b>7XC6G</b>	Genome Sequencing-Parent Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7VRMG</b>	GenomeSeqDx - Duo	GeneDx
<b>7VRNG</b>	GenomeSeqDx - Proband	GeneDx
<b>6DX7G</b>	GenomeSeqDx - Trio	GeneDx
<b>749YG</b>	Genomic Unity Whole Genome Analysis	Variantyx, Inc.
<b>2NVZG</b>	GenomX Sequencing Test	Gene by Gene
<b>86PCG</b>	IriSight? Comprehensive Analysis – Prenatal	Variantyx, Inc.
<b>6R9PG</b>	MNGenome Additional Comparator (only available when trio is ordered)	MNG Laboratories
<b>6R9SG</b>	MNGenome DUO Sequencing	MNG Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6R9QG</b>	MNGenome Proband Only Sequencing	MNG Laboratories
<b>6RLTG</b>	MNGenome TRIO Sequencing	MNG Laboratories
<b>6S6NG</b>	PGnome Diagnostic - Duo	PreventionGenetics, part of Exact Sciences
<b>6S6QG</b>	PGnome Diagnostic - Trio	PreventionGenetics, part of Exact Sciences
<b>4E8TG</b>	PGnome Health Screen Patient Only	PreventionGenetics, part of Exact Sciences
<b>6LTUG</b>	PGnomeTM - Whole Genome Sequencing Patient Only	PreventionGenetics, part of Exact Sciences
<b>7V9UG</b>	Proband WGS	AiLife Diagnostics
<b>86SXG</b>	Proband Whole Genome Sequencing	Baylor Genetics, LLC
<b>7XC8G</b>	Rapid Genome Sequencing - Parent Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7YNNG</b>	Reflex to Whole Genome Sequencing (from genome panels)	Revvity
<b>7RQAG</b>	Standard/DxO Whole Genome Sequencing Additional Family Member	Rady Children's Institute for Genomic Medicine
<b>7RQBG</b>	Standard/DxO Whole Genome Sequencing Proband Only	Rady Children's Institute for Genomic Medicine
<b>7RQCG</b>	Standard/DxO Whole Genome Sequencing Trio	Rady Children's Institute for Genomic Medicine
<b>7V9WG</b>	Trio WGS	AiLife Diagnostics
<b>4ZLBG</b>	Trio Whole Genome Sequencing	Baylor Genetics, LLC
<b>22HVG</b>	TruGenome Proband	Illumina, Inc.
<b>2F7GG</b>	TruGenome Trio	Illumina, Inc.
<b>7U3WG</b>	Whole Genome Sequencing	ARUP Laboratories
<b>7YWTG</b>	Whole Genome Sequencing	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>52SUG</b>	Whole Genome Sequencing	Gene by Gene
<b>7YU6G</b>	Whole Genome Sequencing	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>7Z2QG</b>	Whole Genome Sequencing	Rady Children's Institute for Genomic Medicine
<b>27KEG</b>	Whole Genome Sequencing	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>6RPTG</b>	Whole Genome Sequencing	Children's Mercy Hospital and Clinics - Molecular Genetics Laboratory
<b>7TYTG</b>	Whole Genome Sequencing (Baylor)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7SAVG</b>	Whole Genome Sequencing for Hereditary Disorders, Varies	Mayo Clinic Laboratories

Centene TX Ambetter Superior Genetic Testing and Laboratory Matrix 09/1/2025.docx  
 9/1/2025

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YLSG</b>	Whole Genome Sequencing Proband Only	Revvity
<b>22DHG</b>	Whole Genome Sequencing Single and Multi-Family (Duo, Trio, Quad, etc.)	Medical College of Wisconsin - Human and Molecular Genetics Center
<b>7TKXG</b>	Whole Genome Sequencing, 4 Sample	Praxis Genomics, LLC
<b>7TKVG</b>	Whole Genome Sequencing, DUO	Praxis Genomics, LLC
<b>7YLJG</b>	Whole Genome Sequencing, DUO	Revvity
<b>7U3XG</b>	Whole Genome Sequencing, Familial Control	ARUP Laboratories
<b>7TKUG</b>	Whole Genome Sequencing, Proband	Praxis Genomics, LLC
<b>7YLFG</b>	Whole Genome Sequencing, QUAD	Revvity
<b>7YLMG</b>	Whole Genome Sequencing, TRIO	Revvity
<b>7TKWG</b>	Whole Genome Sequencing, TRIO	Praxis Genomics, LLC
<b>7TLHG</b>	Whole Genome Upgrade per Sample	Praxis Genomics, LLC
<b>7TLNG</b>	Whole Genome Upgrade per Sample	Praxis Genomics, LLC
<b>7YFJG</b>	Whole Genome Sequencing, Carrier Couple	Praxis Genomics, LLC
<b>7YFKG</b>	Whole Genome Sequencing, Carrier Individual	Praxis Genomics, LLC
<b>7YLRG</b>	Healthy Whole Genome Sequencing, proband only	Revvity
<b>7YLAG</b>	Reflex to Whole Genome Sequencing TRIO (from WGS Proband)	Revvity
<b>7YL9G</b>	Whole Genome Sequencing DATA Only (per sample)	Revvity
<b>7YLCG</b>	Whole Genome Sequencing, QUINT	Revvity
<b>7YQAG</b>	Genomic Unity 2.0	Variantyx, Inc.
<b>7XC7G</b>	Genome Sequencing-Patient Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7XC9G</b>	Rapid Genome Sequencing-Patient Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory

### Hematologic Malignancy Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2DVSG</b>	Comprehensive Lymphoid Oncology (DNA and RNA analysis)	Children's Hospital Colorado Precision Diagnostics Laboratory

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>5M87G</b>	FISH Panel : Myeloid Disorders	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>6Z8DG</b>	FoundationOne Heme	Foundation Medicine
<b>7XE7G</b>	Hematologic Cancer Fusion Analysis	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>73H3G</b>	Hematologic Neoplasm Mutation Panel	Ohio State University - Molecular Pathology Laboratory
<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>6LTNG</b>	Heme Gene Panel by NGS	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>72AHG</b>	IntelliGEN Myeloid	LabCorp
<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>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>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
<b>8EFPG</b>	Myeloid Neoplasm Mutation Analysis Only	Medfusion
<b>6UY4G</b>	Myeloid tumor panel - Somatic Mutation Analysis	Centogene
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>86SCG</b>	Tempus  xT: Targeted panel of 648 genes (Hematologic Malignancy)	Tempus AI, Inc.
<b>7RJGG</b>	JMML Associated Exon Panel (JMML)	Rady Children's Institute for Genomic Medicine
<b>7SFFG</b>	Comprehensive HemeComplete Profile	PathGroup
<b>7SFEG</b>	Comprehensive Myeloid Profile	PathGroup
<b>7SFCG</b>	HemeComplete NextGen Sequencing Assay	PathGroup
<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>7SFBG</b>	Myeloid NextGen Sequencing Assay	PathGroup
<b>7SPKG</b>	Neo Comprehensive - Myeloid Disorders	NeoGenomics Laboratories
<b>45EWG</b>	Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7C5AG</b>	OnkoSight Advanced NGS Myeloid Panel	BioReference Laboratories
<b>7T3WG</b>	OnkoSight Advanced Pan Heme Fusion NGS Panel	BioReference Laboratories
<b>7DN8G</b>	Paired Tumor / Normal - Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7DN9G</b>	Paired Tumor / Normal – Comprehensive Hematologic Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7TS6G</b>	Targeted Oncology Panel Next Generation Sequencing Bone Marrow	Cleveland Clinic Laboratories
<b>2FKWG</b>	Comprehensive 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>7UXCG</b>	Myeloid Malignancies Mutation and Copy Number Variation Panel by Next Generation Sequencing	ARUP Laboratories
<b>7URQG</b>	Neo Comprehensive - Heme Cancers	NeoGenomics Laboratories
<b>7VMHG</b>	FISH Panel: Myeloid Malignancy	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7ATHG</b>	HEME DNA/RNA COMBINED PANEL (Combined heme mutation and fusion panels)	Texas Children's Hospital
<b>6LTQG</b>	Myeloid Gene Panel by NGS	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory

GTU	Test Name	Laboratory Name
7ALDG	MYELOID MALIGNANCIES MUTATION PANEL	Clinical Pathology Laboratories
78N9G	Myeloid Multigene Panel	UCSF Molecular Diagnostics Laboratory
7Q2MG	Myeloid Mutation Panel, Blood	Nebraska Medical Center - Molecular Diagnostic Laboratory
7Q2HG	Myeloid Mutation Panel, Other	Nebraska Medical Center - Molecular Diagnostic Laboratory
5CTBG	Myeloid Mutation Panel- AML	University of North Carolina Hospitals - Molecular Genetics
7V4RG	Myeloid NGS	Allina Health Laboratory
7TV4G	Myeloid NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
7SQ8G	Myeloid Panel NGS Bone Marrow	Cleveland Clinic Laboratories
7SQ7G	Myeloid Panel NGS Peripheral Blood	Cleveland Clinic Laboratories
2LJDG	NGS Hematolymphoid Panel (Lab Only)	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
5364G	NGS_Myeloid 37 Genes Panel	CellNetix Pathology and Laboratories
86LPG	OnkoSight Advanced Chronic Lymphoid Neoplasm NGS Panel - PB/BM	BioReference Laboratories
7ZQTG	Targeted Myeloid Panel (TMP)	Columbia University - Personalized Genomic Medicine
7XR9G	MayoComplete Chronic Lymphoid Neoplasms, Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
7XR8G	MayoComplete Histiocytic Neoplasms, Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
7XR7G	MayoComplete T-Cell Lymphoma, Next-Generation Sequencing, Varies	Mayo Clinic Laboratories
34YSG	GeneTrails Hematologic Malignancies 220 Gene Panel	Knight Molecular Diagnostic
7XY8G	MYD88 and CXCR4 Mutation Panel	Knight Molecular Diagnostic
7SEJG	NGS Hematology Molecular Profile	Palo Verde Laboratory - division of Sonora Quest Lab
7W6YG	T Large Granular Lymphocytic Leukemia NGS Panel	University of Minnesota Physicians Outreach Laboratory
7Z2DG	NGS Hematologic Malignancy Mutation Panel (MHIME)	Rady Children's Institute for Genomic Medicine

### Hereditary GI/Colon Cancer Panel Tests

GTU	Test Name	Laboratory Name
2NBHG	CentoColon - NGS Panel (CNV included)	Centogene

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>7WT3G</b>	ColoNext	University of Minnesota Physicians Outreach Laboratory
<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>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>76DWG</b>	Colorectal Cancer Panel	GeneDx
<b>76DSG</b>	Colorectal Cancer Panel	ACL Laboratories
<b>7YQ5G</b>	Colorectal Cancer Panel	Revvity
<b>3DW5G</b>	ColoSeq - Lynch and Polyposis Panel	University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory
<b>6V2QG</b>	Comprehensive Colon Cancer Panel (Seq & Del/Dup)	Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories
<b>76DZG</b>	Hereditary Colorectal Cancer Panel	The University of Chicago Genetic Services
<b>25LZG</b>	Hereditary Colorectal Cancer Panel	Quest Diagnostics
<b>7V2BG</b>	Hereditary Colorectal/Gastrointestinal Cancer Panel	Baylor Genetics, LLC
<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>7V2GG</b>	Hereditary High-Risk Colorectal Cancer Panel	Baylor Genetics, LLC

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>45K7G</b>	Inherited Colon Cancer Panel	Knight Molecular Diagnostic
<b>6DXSG</b>	Lynch/Colorectal High Risk Panel	GeneDx
<b>7YQ4G</b>	STAT Colorectal Cancer Panel	Revvity
<b>76CBG</b>	VistaSeq Colorectal Cancer Panel	Labcorp   Oncology
<b>76DAG</b>	VistaSeq Colorectal Cancer Panel	LabCorp
<b>76BSG</b>	VistaSeq Colorectal Cancer Panel	Integrated Genetics
<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
<b>7SHSG</b>	Hereditary Gastrointestinal Cancer High-Risk Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<b>7TP3G</b>	CancerNext Expanded Colon Lynch	Ambry Genetics
<b>7TNZG</b>	CancerNext: Colon/Lynch	Ambry Genetics
<b>7TNXG</b>	Comprehensive Common Cancer Panel: Colon/Lynch	GeneDx
<b>7TNYG</b>	Rest of Comprehensive Common Cancer Panel: Colon/Lynch	GeneDx
<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>6U88G</b>	Colorectal Cancer Focus Panel (Germline)	NeoGenomics Laboratories
<b>7AAAG</b>	Hereditary Colorectal Cancer Panel (20 Genes)	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7X25G</b>	GxVISION Hereditary Cancer Risk Assessment Colorectal Cancer Genes	Otogenetics
<b>7YHKG</b>	Tempus xG CancerNext 39 genes (hereditary colon cancer indications)	Tempus AI, Inc.
<b>7YHAG</b>	Tempus xG CancerNext 39 genes + RNA (hereditary colon cancer indications)	Tempus AI, Inc.

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YHJG</b>	Tempus xG CancerNext-Expanded 76 genes (hereditary colon cancer indications)	Tempus AI, Inc.
<b>7YHFG</b>	Tempus xG CancerNext-Expanded 76 genes + RNA (hereditary colon cancer indications)	Tempus AI, Inc.
<b>7YQDG</b>	OncoAlly Hereditary Colorectal Cancer Analysis	Variantyx, Inc.
<b>6V3XG</b>	Hereditary Colorectal Cancer and Polyposis Panel	PreventionGenetics, part of Exact Sciences

### Hereditary Polyposis Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>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>7TQFG</b>	Adenomatous Polyposis Focus (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7TQGG</b>	Polyposis Comprehensive (Sequencing & Deletion/Duplication)	Fulgent Genetics

### HLA-DQ Typing (Celiac Disease) Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2A2PG</b>	Celiac Associated HLA-DQ genotyping	Cleveland Clinic Laboratories
<b>5NUVG</b>	Celiac Disease (HLA-DQ2 and HLA-DQ8) Genotyping (2005018) (Back-Up Only)	Rady Children's Institute for Genomic Medicine
<b>7BGEG</b>	CELIAC DISEASE GENOTYPE	Clinical Pathology Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>3HMZG</b>	Celiac Disease HLA DQ Assoc	Integrated Regional Laboratories
<b>483JG</b>	Celiac Disease HLA DQ association	Allina Health Laboratory
<b>7XM9G</b>	Celiac Disease HLA Screen (OUTREACH)	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>4PZ4G</b>	Celiac Disease HLA Screen (UIMC)	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>4H24G</b>	Celiac Disease HLA Typing	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>7DJXG</b>	Celiac Disease HLA-DQ Genotyping	ARUP Laboratories
<b>4PSLG</b>	Celiac Genetics	Norton CPA Lab
<b>4N8CG</b>	Celiac Genetics	Children's Mercy Hospital and Clinics - Molecular Genetics Laboratory
<b>7Y4LG</b>	Celiac HLA DQ Association	LabCorp
<b>7Y4FG</b>	Celiac HLA DQ Association with Reflex to Celiac Antibodies tTG IgA, tTG IgG, DGP IgA, DGP IgG and Total IgA	LabCorp
<b>6ZNDG</b>	HLA Celiac	UCSF Molecular Diagnostics Laboratory
<b>7UE8G</b>	HLA Celiac	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>5XHBG</b>	HLA Celiac - DQA1, DQB1	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>38Q5G</b>	HLA Celiac Disease Association (HLA-DQ2/DQ8)	University of Michigan - Michigan Medical Genetics Laboratories
<b>3HN2G</b>	HLA DQ2/DQ8 for Celiac Disease	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>8EGHG</b>	HLA Typing for Celiac Disease	Medfusion
<b>3A69G</b>	HLA Typing for Celiac Disease (17135)	Rady Children's Institute for Genomic Medicine
<b>4QTYG</b>	HLA-DQ Genotyping	ARUP Laboratories
<b>38QUG</b>	HLA-DQ2/DQ8/DQA105 (Celiac association)	Bloodworks Northwest - Hemostasis, Genomics, and Platelet Lab

#### HPV-Related Solid Tumor Cell-free DNA (cfDNA) Panel Tests for Monitoring Minimal Residual Disease (MRD)

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YF4G</b>	HPV-SEQ	Sysmex Inostics, INC
<b>7VX2G</b>	NavDx	Naveris

## Hypertrophic Cardiomyopathy (HCM) Panel Tests

GTU	Test Name	Laboratory Name
<b>2YZAG</b>	HCMNext Reflex	Ambry Genetics
<b>2YZEG</b>	HCMNext	Ambry Genetics
<b>3CEJG</b>	Hypertrophic Cardiomyopathy (HCM) Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7DN2G</b>	Hypertrophic Cardiomyopathy (HCM) Panel	Washington University in St. Louis Genomics and Pathology Services
<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>7XH6G</b>	Hypertrophic Cardiomyopathy Panel	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7UVVG</b>	Hypertrophic Cardiomyopathy Panel	The University of Chicago Genetic Services
<b>3XQ7G</b>	Hypertrophic Cardiomyopathy Panel	Knight Molecular Diagnostic
<b>5CYCG</b>	Hypertrophic Cardiomyopathy Panel	PreventionGenetics, part of Exact Sciences
<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>42LUG</b>	Invitae Hypertrophic Cardiomyopathy Panel-Add-on Preliminary-evidence Genes for Hypertrophic Cardiomyopathy	Invitae Corporation
<b>7SBZG</b>	Hypertrophic Cardiomyopathy Gene Panel, Varies	Mayo Clinic Laboratories
<b>7YWCG</b>	Invitae Hypertrophic Cardiomyopathy Panel - ROC01	Invitae Corporation
<b>7YWBG</b>	Invitae Hypertrophic Cardiomyopathy Panel-Add-on Preliminary-evidence Genes for Hypertrophic Cardiomyopathy - ROC01	Invitae Corporation

## Long QT Syndrome (LQTS) Panel Tests

GTU	Test Name	Laboratory Name
<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>2KQNG</b>	Long QT Panel, Sequencing and Deletion/Duplication	ARUP Laboratories
<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>7VUUG</b>	Long QT Syndrome Panel	The University of Chicago Genetic Services
<b>7XHMG</b>	Long QT Syndrome Panel	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>76V5G</b>	Long QT Syndrome Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>78FTG</b>	Long QT Syndrome Panel	PreventionGenetics, part of Exact Sciences
<b>7YK8G</b>	Long QT Syndrome Panel	Revvity
<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>7YK7G</b>	STAT Long QT Syndrome Panel	Revvity
<b>7SBNG</b>	Long QT Syndrome Gene Panel, Varies	Mayo Clinic Laboratories

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

GTU	Test Name	Laboratory Name
<b>6RPMG</b>	COLARIS	Myriad Genetics
<b>2YYWG</b>	HNPCC concurrent	Ambry Genetics
<b>2FTJG</b>	HNPCC PANEL	Center for Human Genetics, Inc.
<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>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>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>2YYNG</b>	MSH2 seq & del/dup & EPCAM del/dup	Ambry Genetics
<b>87B9G</b>	VistaSeq Lynch Syndrome Panel	LabCorp
<b>7YJ3G</b>	MLH1/MSH2 Del/Dup Testing by MLPA	Revvity
<b>7YJ4G</b>	MSH6/MUTYH/EPCAM Del/Dup Testing by MLPA	Revvity
<b>7QFCG</b>	Lynch Syndrome Focus (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>7SNVG</b>	HNPCC/Lynch Deletion/Duplication	Ambry Genetics
<b>7U2HG</b>	MLH1, MSH2, MSH6, PMS2 Sequencing and Del/Dup (NGS)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7X23G</b>	GxVISION Hereditary Cancer Risk Assessment Lynch Syndrome Genes	Otogenetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>57DNG</b>	Lynch Syndrome/Constitutional Mismatch Repair Deficiency Panel	PreventionGenetics, part of Exact Sciences

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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>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>7YP7G</b>	Comprehensive Mitochondrial Nuclear Gene Panel	Revvity
<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, Inc.
<b>7V37G</b>	Genomic Unity Mitochondrial Genome Deletions Analysis	Variantyx, Inc.
<b>7V36G</b>	Genomic Unity Nuclear Encoded Mitochondrial Gene Analysis	Variantyx, 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>77GLG</b>	Mitochondrial Complex I Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<b>6QVJG</b>	Mitochondrial Complex II Deficiency Panel	PreventionGenetics, part of Exact Sciences
<b>386QG</b>	Mitochondrial Complex III Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76ZNG</b>	Mitochondrial Complex IV Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<b>5LQTG</b>	Mitochondrial Complex V Deficiency Panel (Nuclear Genes)	PreventionGenetics, part of Exact Sciences
<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>54EFG</b>	Mitochondrial Encephalomyopathic Evaluation	Athena Diagnostics Inc
<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>77T2G</b>	Mitochondrial Respiratory Chain Complex I-V Nuclear Gene Deficiency Panel by Massively Parallel Sequencing	Baylor Genetics, LLC
<b>4RDTG</b>	Mitochondrial Respiratory Chain Complex II 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>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>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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>7YP6G</b>	STAT Comprehensive Mitochondrial Nuclear Gene Panel	Revuity
<b>6XJWG</b>	Mitochondrial DNA Depletion Testing (Leukocyte)	LabCorp

### 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>7V38G</b>	Genomic Unity Mitochondrial Genome Sequence Analysis	Variantyx, Inc.
<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>253UG</b>	Mitochondrial Disorders (mtDNA) Sequencing and Deletion Analysis by NGS	ARUP Laboratories

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>72G9G</b>	Mitochondrial DNA Depletion Testing (Muscle)	LabCorp
<b>5D36G</b>	Mitochondrial DNA Sequencing	New Jersey Medical School - Institute of Genomic Medicine
<b>6R5TG</b>	Mitochondrial Full Genome Analysis, Next-Generation Sequencing (NGS), Varies	Mayo Clinic Laboratories
<b>6R9BG</b>	MITOCHONDRIAL GENOME SEQUENCING	MNG Laboratories
<b>7RXFG</b>	MITOCHONDRIAL GENOME SEQUENCING	LabCorp
<b>4CH3G</b>	MITOCHONDRIAL GENOME SEQUENCING	Center for Human Genetics, Inc.
<b>6R9DG</b>	Mitochondrial Genome Sequencing + Deletion Analysis	MNG Laboratories
<b>7RXEG</b>	Mitochondrial Genome Sequencing and Deletion Analysis	LabCorp
<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
<b>4KWSG</b>	Mitochondrial Whole Genome Sequence Analysis (Familial Mutation/Variant Analysis)	Baylor Genetics, LLC
<b>7Z28G</b>	MT DNA Sequencing and Deletion Analysis (554)	Rady Children's Institute for Genomic Medicine

#### Noonan Spectrum Disorders/RASopathies Panel Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>3XMZG</b>	11-GENE PANEL FOR PTPN11;SOS1;KRAS;SHOC2;NRAS; RAF1; BRAF, CBL, SOS2; RIT1; AND LZTR1	Center for Human Genetics, Inc.
<b>3EBYG</b>	4-GENE PANEL FOR PTPN11;SOS1;KRAS; SHOC2	Center for Human Genetics, Inc.
<b>3YXGG</b>	6-GENE PANEL FOR PTPN11;SOS1;KRAS;SHOC2;NRAS; RAF1	Center for Human Genetics, Inc.
<b>6U7MG</b>	Expanded NF1 RASopathy Panel by Next-Generation Sequencing and Deletion/Duplication Analysis of LZTR1, NF1, and SPRED1 (RAS-NG)	Department of Genetics UAB - Medical Genomics Laboratory
<b>56ZDG</b>	GeneSeq Cardio: Noonan Syndrome / RASopathies Panel	LabCorp

<b>2Y77G</b>	GeneSeq: Cardio-Noonan Syndrome / RASopathies Profile	Integrated Genetics
<b>2AXLG</b>	Invitae RASopathies and Noonan Spectrum Disorders Panel	Invitae Corporation
<b>378GG</b>	KRAS/RAF1/SOS1 DNA Sequencing Evaluation	Athena Diagnostics Inc
<b>3D3YG</b>	LEOPARD SYNDROME PANEL	Center for Human Genetics, Inc.
<b>6RACG</b>	Non-NF1 RASopathy Panel by Next-Generation Sequencing and Deletion/Duplication Analysis of LZTR1 and SPRED1 (NNP-NG)	Department of Genetics UAB - Medical Genomics Laboratory
<b>79Q8G</b>	Noonan - RASopathies panel - NGS Panel (CNV included)	Centogene
<b>6DSVG</b>	Noonan and Comprehensive RASopathies Panel	GeneDx
<b>77YJG</b>	Noonan and Comprehensive RASopathies Panel	Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital
<b>475FG</b>	Noonan and RASopathies NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<b>25DTG</b>	Noonan and RASopathies NGS Panel (Sequencing & Deletion/Duplication)	Fulgent Genetics
<b>52VKG</b>	Noonan and RASopathies NGS Panel (Sequencing Only)	Fulgent Genetics
<b>2Z83G</b>	Noonan Disorder Gene Panel	Ann and Robert Lurie Children's Hospital of Chicago
<b>28VNG</b>	Noonan Disorders Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>46HVG</b>	Noonan Spectrum Disorder Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>255AG</b>	Noonan Spectrum Disorders Panel, Sequencing	ARUP Laboratories
<b>257XG</b>	Noonan Spectrum Disorders Panel, Sequencing, Fetal	ARUP Laboratories
<b>86CFG</b>	Noonan Spectrum Disorders/RASopathies Panel	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>76J2G</b>	Noonan Spectrum Disorders/RASopathies Panel	PreventionGenetics, part of Exact Sciences
<b>7V2DG</b>	Noonan Spectrum Disorders/RASopathy Panel	Baylor Genetics, LLC
<b>2SCBG</b>	Noonan Syndrome	Norton CPA Lab
<b>2BYKG</b>	Noonan Syndrome (NGS Panel and Copy Number Analysis)	MNG Laboratories
<b>7SBBG</b>	Noonan Syndrome and Related Conditions Gene Panel, Varies	Mayo Clinic Laboratories
<b>5NFZG</b>	Noonan Syndrome Panel	The University of Chicago Genetic Services
<b>2L2VG</b>	Noonan Syndrome Tiered Testing	University of Michigan - Michigan Medical Genetics Laboratories
<b>2YY8G</b>	NoonanNext	Ambry Genetics
<b>6L7JG</b>	Prenatal Noonan Spectrum Disorders Panel	GeneDx
<b>7VXNG</b>	Prenatal Noonan Spectrum Disorders/RASopathy Panel	Baylor Genetics, LLC

Centene TX Ambetter Superior Genetic Testing and Laboratory Matrix 09/1/2025.docx  
 9/1/2025

<b>4ZP7G</b>	Prenatal Noonan Syndrome	Integrated Genetics
<b>7VV8G</b>	RASopathies and Noonan Spectrum Disorders Panel	The University of Chicago Genetic Services
<b>7Y2FG</b>	Rasopathies Panel	Washington University in St. Louis Genomics and Pathology Services
<b>5N8UG</b>	RASopathy NGS Panel	Greenwood Genetic Center - Molecular Diagnostic Laboratory
<b>78GWG</b>	RASopathy/Noonan Spectrum Disorders Panel	Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories
<b>7VBPG</b>	RASopathy/Noonan Syndrome NGS Panel	AiLife Diagnostics
<b>7VBNG</b>	RASopathy/Noonan Syndrome NGS Panel Rapid	AiLife Diagnostics

#### Ovarian Cancer Treatment Algorithmic Tests

GTU	Test Name	Laboratory Name
<b>6RN6G</b>	myChoice CDx	Myriad Genetics
<b>7TV6G</b>	Myriad myChoice CDx	University of Michigan - Michigan Medical Genetics Laboratories

#### Pan-Cancer Hereditary Cancer Panel Tests

GTU	Test Name	Laboratory Name
<b>76DGG</b>	CancerNext	Ambry Genetics
<b>76E2G</b>	CancerNext-Expanded	Ambry Genetics
<b>6UR7G</b>	CancerNext-Expanded with RNAinsight	Ambry Genetics
<b>6UR6G</b>	CancerNext 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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76BTG</b>	COMP CANCER PANEL	ACL Laboratories
<b>7YPZG</b>	Comprehensive Cancer Panel	Revvity
<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>262EG</b>	Comprehensive Hereditary Cancer Panel	Quest Diagnostics
<b>7BP4G</b>	Comprehensive Hereditary Cancer Panel	The University of Chicago Genetic Services
<b>5K7ZG</b>	Comprehensive Hereditary Cancer Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>76DVG</b>	Comprehensive Hereditary Cancer Panel	PreventionGenetics, part of Exact Sciences
<b>7V28G</b>	Comprehensive Hereditary Cancer Panel	Baylor Genetics, LLC
<b>76DHG</b>	CustomNext-Cancer	Ambry Genetics
<b>7WJYG</b>	CustomNext-Cancer	University of Minnesota Physicians Outreach Laboratory
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76DRG</b>	Hereditary Cancer	Color Genomics
<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>76DKG</b>	Inherited Cancer Panel	Knight Molecular Diagnostic
<b>76CHG</b>	Integrated BRACAnalysis with Myriad myRisk Hereditary Cancer Update Test	Myriad Genetics
<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>76CWG</b>	OncoGene Dx Custom Panel	GeneDx
<b>2TQXG</b>	PrevenTest	Advanced Molecular Diagnostics, LLC
<b>7YPYG</b>	STAT Comprehensive Cancer Panel	Revuity
<b>76BRG</b>	VistaSeq Hereditary Cancer Panel	Labcorp   Oncology
<b>76BXG</b>	VistaSeq Hereditary Cancer Panel	LabCorp
<b>76D8G</b>	VistaSeq Hereditary Cancer Panel	Integrated Genetics
<b>7DQGG</b>	NxGen Hereditary Cancer Panel	NxGen MDx
<b>7RVTG</b>	Riskguard	Exact Sciences Laboratories, LLC
<b>7YHXG</b>	WholeCancerTM Panel	Revuity
<b>7RX8G</b>	Common Hereditary Cancer Screening Panel	PreventionGenetics, part of Exact Sciences
<b>7TP4G</b>	CancerNext Expanded HBOC	Ambry Genetics
<b>7TP2G</b>	CancerNext: HBOC	Ambry Genetics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>76C3G</b>	Comprehensive Common Cancer Panel: HBOC	GeneDx
<b>7VAGG</b>	Hereditary Cancer NGS Panel	AiLife Diagnostics
<b>7VAHG</b>	Hereditary Cancer NGS Panel Rapid	AiLife Diagnostics
<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>7VZFG</b>	GeneticsNow Comprehensive	GoPath Laboratories
<b>7X24G</b>	GxVISION Hereditary Cancer Risk Assessment Comprehensive Inherited Cancer Gene Tests	Otogenetics
<b>7XZNG</b>	Oncopanel germline: custom capture next generation sequencing for cancer risk	Brigham and Women's Hospital - Center for Advanced Molecular Diagnostics
<b>7AABG</b>	Comprehensive Hereditary Cancer Panel (66 Genes)	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7AA9G</b>	Guideline Based Hereditary Cancer Pnl (32 Genes)	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7YHBG</b>	Tempus xG CancerNext 39 genes (all other hereditary cancer indications)	Tempus AI, Inc.
<b>7YH8G</b>	Tempus xG CancerNext 39 genes + RNA (all other hereditary cancer indications)	Tempus AI, Inc.
<b>7YHGG</b>	Tempus xG CancerNext-Expanded 76 genes (all other hereditary cancer indications)	Tempus AI, Inc.
<b>7YHDG</b>	Tempus xG CancerNext-Expanded 76 genes + RNA (all other hereditary cancer indications)	Tempus AI, Inc.
<b>7YT9G</b>	FoundationOneGermline	Foundation Medicine
<b>7YTAG</b>	FoundationOneGermline More	Foundation Medicine

### 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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2AUXG</b>	GeneSight Psychotropic	Myriad Genetics
<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>7YC9G</b>	Pharmacogenetics Panel: Psychotropics	University of Illinois at Chicago - Biochemical Genetics Laboratory
<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>33HHG</b>	Tempus nP	Tempus AI, Inc.
<b>7WZVG</b>	Otogenetics PGx Mental Health	Otogenetics
<b>7Y2XG</b>	iPsychGx	iGenomeDx

### Prenatal Cell-Free RH DNA Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7XQCG</b>	Fetal RhD NIPT Add-On	Natera
<b>7YSTG</b>	PrenatalDetect RHD	Devyser Genomic Laboratories
<b>7T56G</b>	UNITY Fetal RhD™ NIPT (add on)	BillionToOne, Inc.

### Prostate Cancer Treatment and Prognostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UUUG</b>	Polaris	Myriad Genetics
<b>6UU7G</b>	Polaris Biopsy	Myriad Genetics
<b>7BN5G</b>	Decipher Prostate Genomic Classifier	Veracyte
<b>7Q8UG</b>	Genomic Prostate Score Test	MDx Health

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7Y3JG</b>	ArteraAI	Artera
<b>7YSUG</b>	PROSTOX	MiraDx

### Rapid Exome Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7U23G</b>	Critical Trio Whole Exome Sequencing and Comprehensive mtDNA Analysis	University of Michigan - Michigan Medical Genetics Laboratories
<b>2ZYHG</b>	ExomeNext-Rapid	Ambry Genetics
<b>6RLSG</b>	MNG STAT Exome DUO Sequencing	MNG Laboratories
<b>33PKG</b>	MNG STAT Exome DUO Sequencing + mtDNA	MNG Laboratories
<b>6RLQG</b>	MNG STAT Exome Proband Only Sequencing	MNG Laboratories
<b>49JPG</b>	MNG STAT Exome Proband Only Sequencing + mtDNA	MNG Laboratories
<b>6RGLG</b>	MNG STAT Exome TRIO Sequencing	MNG Laboratories
<b>6RLGG</b>	MNG STAT Exome TRIO Sequencing + mtDNA	MNG Laboratories
<b>25EZG</b>	PGxome RAPID Exome Test - Patient Only	PreventionGenetics, part of Exact Sciences
<b>25GEG</b>	PGxome RAPID Exome Test - Trio	PreventionGenetics, part of Exact Sciences
<b>6U84G</b>	PGxome RAPID Exome Test Family Duo	PreventionGenetics, part of Exact Sciences
<b>7VBZG</b>	Rapid Duo Exome	AiLife Diagnostics
<b>86SVG</b>	Rapid Duo Whole Exome Sequencing	Baylor Genetics, LLC
<b>7VBVG</b>	Rapid Proband Exome	AiLife Diagnostics
<b>7R8DG</b>	Rapid Proband Whole Exome Sequencing	Baylor Genetics, LLC
<b>7VBXG</b>	Rapid Trio Exome	AiLife Diagnostics
<b>25EFG</b>	Rapid Trio Whole Exome Sequencing	Baylor Genetics, LLC
<b>25FKG</b>	Rapid Trio Whole Exome Sequencing + Comprehensive mtDNA Analysis	Baylor Genetics, LLC
<b>7YM2G</b>	Rapid Whole Exome Sequencing DUO	Revvity
<b>7YM8G</b>	Rapid Whole Exome Sequencing Proband Only	Revity
<b>7YLXG</b>	Rapid Whole Exome Sequencing QUAD	Revity

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YM6G</b>	Rapid Whole Exome Sequencing TRIO	Revvity
<b>7YLVG</b>	Rapid Whole Exome Sequencing, Quint	Revvity
<b>25ENG</b>	STAT Exome Sequencing (Proband Only)	The University of Chicago Genetic Services
<b>25EJG</b>	STAT Exome Sequencing (Trio)	The University of Chicago Genetic Services
<b>7YM3G</b>	STAT Prenatal Whole Exome Sequencing, DUO	Revvity
<b>7YM9G</b>	STAT Prenatal Whole Exome Sequencing, Proband ONLY	Revvity
<b>7LYLG</b>	STAT Prenatal Whole Exome Sequencing, QUAD	Revvity
<b>7YM5G</b>	STAT Prenatal Whole Exome Sequencing, TRIO	Revvity
<b>25EXG</b>	WES Trio Rapid	Medical College of Wisconsin - Human and Molecular Genetics Center
<b>7VREG</b>	XomeDxXpress - Duo - Rapid Exome Sequencing	GeneDx
<b>7VRPG</b>	XomeDxXpress - Proband - Rapid Exome Sequencing	GeneDx
<b>5MK2G</b>	XomeDxXpress - Trio - Rapid Exome Sequencing	GeneDx

## Rapid Genome Sequencing Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6R9TG</b>	MNGenome STAT DUO Sequencing	MNG Laboratories
<b>6R9RG</b>	MNGenome STAT Proband Only Sequencing	MNG Laboratories
<b>6R9NG</b>	MNGenome STAT TRIO Sequencing	MNG Laboratories
<b>3VAEG</b>	PGnome Rapid	PreventionGenetics, part of Exact Sciences
<b>3CSHG</b>	PGnome - RAPID - Duo	PreventionGenetics, part of Exact Sciences
<b>3VD3G</b>	PGnome - RAPID - Trio	PreventionGenetics, part of Exact Sciences
<b>7V9ZG</b>	Rapid Duo WGS	AiLife Diagnostics
<b>86SYG</b>	Rapid Duo Whole Genome Sequencing	Baylor Genetics, LLC
<b>7V9VG</b>	Rapid Proband WGS	AiLife Diagnostics
<b>86SZG</b>	Rapid Proband Whole Genome Sequencing	Baylor Genetics, LLC
<b>7U3JG</b>	Rapid Targeted Analysis of Family Member	Children's Hospital of Philadelphia - Division of Genomic Diagnostics

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7V9XG</b>	Rapid Trio WGS	AiLife Diagnostics
<b>7AUWG</b>	Rapid Trio Whole Genome Sequencing	Baylor Genetics, LLC
<b>7T4JG</b>	Rapid Whole Genome Sequencing	ARUP Laboratories
<b>7RNXG</b>	Rapid Whole Genome Sequencing Duo	Rady Children's Institute for Genomic Medicine
<b>7RNYG</b>	Rapid Whole Genome Sequencing Proband Only	Rady Children's Institute for Genomic Medicine
<b>7RNZG</b>	Rapid Whole Genome Sequencing Trio	Rady Children's Institute for Genomic Medicine
<b>7T4LG</b>	Rapid Whole Genome Sequencing, Familial Control	ARUP Laboratories
<b>7T4KG</b>	Rapid Whole Genome Sequencing, Familial Control with Report	ARUP Laboratories
<b>7YLHG</b>	STAT Prenatal Whole Genome Sequencing, DUO	Revvity
<b>7YLPG</b>	STAT Prenatal Whole Genome Sequencing, Proband ONLY	Revvity
<b>7YLEG</b>	STAT Prenatal Whole Genome Sequencing, QUAD	Revvity
<b>7YLKG</b>	STAT Prenatal Whole Genome Sequencing, TRIO	Revvity
<b>7RSEG</b>	Ultra-rapid Whole Genome Sequencing	Rady Children's Institute for Genomic Medicine
<b>7U24G</b>	Whole Genome Sequencing, Rapid-Duo (Baylor)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7U25G</b>	Whole Genome Sequencing, Rapid-Proband (Baylor)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7VRKG</b>	GenomeXpress - Duo - Rapid Genome Sequencing	GeneDx
<b>7VRLG</b>	GenomeXpress - Proband - Rapid Genome Sequencing	GeneDx
<b>6UNRG</b>	GenomeXpress - Trio - Rapid Genome Sequencing	GeneDx
<b>7YGCG</b>	GenomeXpress, Trio	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7YLNG</b>	Healthy UltraRapid Whole Genome Sequencing, proband only	Revvity
<b>7YLGG</b>	Ultrarapid Whole Genome Sequencing DUO	Revvity
<b>7YLDG</b>	Ultrarapid Whole Genome Sequencing QUAD	Revvity
<b>7YLLG</b>	Ultrarapid Whole Genome Sequencing TRIO	Revvity
<b>7YLQG</b>	Ultrarapid Whole Genome Sequencing, Proband ONLY	Revvity
<b>7YLBG</b>	Ultrarapid Whole Genome Sequencing, QUINT	Revvity
<b>7YQGG</b>	Genomic Unity Lightning Genome Analysis – Neonatal	Variantyx, Inc.
<b>7YQHG</b>	Genomic Unity Lightning Genome Analysis – Pediatric	Variantyx, Inc.

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7YQJG</b>	Genomic Unity Lightning Genome Analysis – Standard	Variantyx, Inc.
<b>7YT8G</b>	GeneDx ultraRapid Genome Sequencing - Proband	GeneDx
<b>7YT5G</b>	Ultra Rapid Duo WGS	AiLife Diagnostics
<b>7YT7G</b>	Ultra Rapid Proband WGS	AiLife Diagnostics
<b>7YT6G</b>	Ultra Rapid Trio WGS	AiLife Diagnostics
<b>7Z2UG</b>	Ultra-Rapid Genome Sequencing – Parent Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory
<b>7Z2TG</b>	Ultra-Rapid Genome Sequencing – Patient Sample	Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory

### 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>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	Molecular and Genomic Pathology Laboratory, University of Pittsburgh Medical Center (UPMC)
<b>7S8XG</b>	Afirma Xpression Atlas	Veracyte

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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>2YTJG</b>	BCR / ABL - t(9;22) by RT-PCR (Quantitative)	Labcorp   Oncology
<b>3LU5G</b>	BCR ABL P210 QUANT	SUNY Upstate Medical University
<b>5PH3G</b>	BCR-ABL Diagnostic, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>4GZ7G</b>	BCR-ABL Diagnostic, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>5W36G</b>	BCR-ABL Mutation	University of Illinois at Chicago - Biochemical Genetics Laboratory

GTU	Test Name	Laboratory Name
<b>5PGYG</b>	BCR-ABL p190 Followup, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>4GLHG</b>	BCR-ABL p190 Followup, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>4AGMG</b>	BCR-ABL p210 Followup, Blood	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>3AUMG</b>	BCR-ABL p210 Followup, Bone Marrow	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>723HG</b>	BCR-ABL1 Gene Rearrangement, Quantitative, PCR	Quest Diagnostics
<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>7BQPG</b>	BCR/ABL by FISH	Allina Health Laboratory
<b>6ND3G</b>	BCR/ABL D-FISH	Center for Genetic Testing at Saint Francis
<b>3KP8G</b>	BCR/ABL qRT PCR, Major p210 and Minor p190 Mutations, Minimal Residual Disease	Molecular Pathology Laboratory Network
<b>2V82G</b>	BCR/ABL Quantification by RT-PCR	BioReference Laboratories
<b>6YXMG</b>	BCR/ABL t(9;22) major (p210) IS Quantitative	Molecular Diagnostic Laboratory - Barnes Jewish Hospital
<b>6LE9G</b>	BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
<b>6UV4G</b>	BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
<b>5SUVG</b>	BCR/ABL1 p190 Quantitative PCR Blood	Cleveland Clinic Laboratories

GTU	Test Name	Laboratory Name
<b>5UA8G</b>	BCR/ABL1 p190 Quantitative PCR Bone Marrow	Cleveland Clinic Laboratories
<b>7TH5G</b>	BCR/ABL1 p210 and p190 Diagnostic PCR Blood	Cleveland Clinic Laboratories
<b>7TH6G</b>	BCR/ABL1 p210 and p190 Diagnostic PCR Bone Marrow	Cleveland Clinic Laboratories
<b>5UAG</b>	BCR/ABL1 p210 Quantitative PCR Blood	Cleveland Clinic Laboratories
<b>6LQRG</b>	BCR/ABL1 p210 Quantitative PCR Bone Marrow	Cleveland Clinic Laboratories
<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 - UCDMC - Molecular and Cytogenetic Laboratory
<b>7BDAG</b>	BCR-ABL1, QUANT, MINOR (p190)	Clinical Pathology Laboratories
<b>7W5EG</b>	BCR ABL1 Major Breakpoint Quant P210	University of Minnesota Physicians Outreach Laboratory
<b>7WVTG</b>	BCR-ABL1 High Sensitivity Major p210	University of Minnesota Physicians Outreach Laboratory
<b>7YHNG</b>	BCR/ABL1 Qualitative Diagnostic Assay with Reflex to BCR/ABL1 p190 Quantitative Assay or BCR/ABL1 p210 Quantitative Assay, Varies (BCRFX)	Marshfield Labs

### Tumor Specific 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>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>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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>3ZETG</b>	Lung Cancer Panel by Next Generation Sequencing	Molecular Pathology Laboratory Network
<b>7R8HG</b>	LungSEQ Concurrent Panel, without PD-L1	Quest Diagnostics
<b>7AUEG</b>	LUNGSEQ Panel	Quest Diagnostics
<b>8EEXG</b>	LUNGSEQ Panel	Medfusion
<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>7BHXG</b>	NeoTYPE DNA & RNA - Lung	NeoGenomics Laboratories
<b>6YZRG</b>	NeoTYPE Lung Tumor Profile	NeoGenomics Laboratories
<b>86LVG</b>	OnkoSight Advanced Comprehensive Lung	BioReference Laboratories
<b>86LLG</b>	OnkoSight Advanced Lung Cancer NGS Panel	BioReference Laboratories
<b>26QHG</b>	Targeted Gene Panel with Fusions, Lung Cancer	Palo Verde Laboratory - division of Sonora Quest Lab
<b>5XBBG</b>	EGFR and KRAS, if both neg, reflex ALK, if neg reflex ROS1	BioReference Laboratories
<b>6U25G</b>	OnkoSight Advanced Lung Cancer Panel, ALK and ROS1 by FISH	BioReference Laboratories
<b>6ST7G</b>	Oncology FISH Analysis - Non-small Cell Lung Carcinoma Panel	Baylor Genetics, LLC
<b>7VZQG</b>	Lung Cancer Mutation Panel	ARUP Laboratories
<b>7AFUG</b>	LUNG CANCER TARGETED GENE	Clinical Pathology Laboratories
<b>7Y3UG</b>	ALK, RET, ROS1 fusion, MET amplification by FISH	Molecular and Genomic Pathology Laboratory, University of Pittsburgh Medical Center (UPMC)
<b>7YF8G</b>	Lung HDPCR	Protean Biodiagnostics

#### 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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>77HHG</b>	Comprehensive Pan-cancer analysis (DNA and RNA analysis)	Children's Hospital Colorado Precision Diagnostics 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>6R4AG</b>	FoundationOne CDx	Foundation Medicine
<b>3V7ZG</b>	GatorSeq (NGS Panel)	University of Florida - Health Pathology Laboratories
<b>86PJM</b>	Guardant360 TissueNext	Guardant Health
<b>7AWEG</b>	MayoComplete Solid Tumor Panel, Next-Generation Sequencing, Tumor	Mayo Clinic Laboratories
<b>6UUMG</b>	MSK-IMPACT	Memorial Sloan Kettering Cancer Center
<b>34WBG</b>	NGS Solid Tumor Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<b>2YA6G</b>	OncoGxOne	Admera Health
<b>76ETG</b>	Oncoplex Select Cancer Gene Panel	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>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>78FJG</b>	Pan-cancer DNA Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>2K9WG</b>	PGDx elio tissue complete	Personal Genome Diagnostics
<b>7A5HG</b>	Precise Tumor	Myriad Genetics
<b>2D5BG</b>	Solid Tumor DNA Analysis	Children's Hospital Colorado Precision Diagnostics Laboratory
<b>3CSSG</b>	Solid Tumor Molecular Profile	Fulgent Genetics
<b>2NSPG</b>	Solid Tumor NGS Panel	University of Michigan - Michigan Medical Genetics Laboratories
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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
<b>6WUTG</b>	MI Tumor Seek Hybrid	Caris Life Sciences
<b>7SVMG</b>	Solid Tumor Expanded Panel	Quest Diagnostics
<b>7VQWG</b>	Comprehensive Solid Tumor Panel	Children's Hospital of Philadelphia - Division of Genomic Diagnostics
<b>7V34G</b>	OncoAlly Solid Tumor Analysis	Variantyx, Inc.
<b>2FRPG</b>	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>7UJGG</b>	Fusion Transcript Panel	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>7UJFG</b>	PennSeq Solid Tumor Panel	University of Pennsylvania School of Medicine - Molecular Pathology Laboratory
<b>7UQMG</b>	Solid TumorSEQ Expanded Panel	Medfusion
<b>2CPUG</b>	Strata Select	Strata Oncology
<b>7SSDG</b>	Neo Comprehensive - Solid Tumor	NeoGenomics Laboratories
<b>7UUYG</b>	OnkoSight Advanced 523 Gene NGS Panel	BioReference Laboratories
<b>7URPG</b>	OnkoSight Advanced 523 Gene NGS with PierianDx Interpretation	BioReference 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>6ZQMG</b>	Iowa Cancer Mutation and RNA Fusion Profile and Interpretation	University of Iowa Hospitals and Clinics - Department of Pathology
<b>7529G</b>	NeXT Dx	Personalis
<b>7SSVG</b>	OnkoSight Advanced 523 Gene NGS Reanalysis with PierianDx	BioReference Laboratories
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>7TXJG</b>	Tempus xT Targeted Panel (Tumor plus Saliva)	University of Michigan - Michigan Medical Genetics Laboratories
<b>7ATMG</b>	SOLID TUMOR COMPREHENSIVE PANEL (Combined solid tumor mutation and fusion panels)	Texas Children's Hospital
<b>7VPEG</b>	Tempus xT CDx: FDA-approved Companion Diagnostic	Tempus AI, Inc.
<b>7BNDG</b>	OncоЕxTra	Exact Sciences Laboratories, LLC
<b>7XMNG</b>	GeneAssure Solid Tumor NGS Assay	University of Illinois at Chicago - Biochemical Genetics Laboratory
<b>292AG</b>	MI Cancer Seek	Caris Life Sciences
<b>7XZMG</b>	Oncopanel: 447 gene custom capture next generation sequencing	Brigham and Women's Hospital - Center for Advanced Molecular Diagnostics
<b>7Y3KG</b>	UPMC Oncomine	Molecular and Genomic Pathology Laboratory, University of Pittsburgh Medical Center (UPMC)
<b>7YEXG</b>	Liquid Trace: Solid Tumor Profile	Sysmex Inostics, INC
<b>3AE8G</b>	Targeted Gene Panel with Fusions, Comprehensive Tumor	Palo Verde Laboratory - division of Sonora Quest Lab
<b>7YSVG</b>	TruSight Oncology Comprehensive	Illumina, Inc.
<b>7YY6G</b>	Iowa Pan-Cancer Mutation Profiling	University of Iowa Hospitals and Clinics - Department of Pathology

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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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>7BZJG</b>	GIST targeted NGS panel	Allina Health Laboratory
<b>5BRWG</b>	NGS Solid Tumor Hotspot Panel	Johns Hopkins Medical Institutions - Pathology Laboratory
<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>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<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
<b>7XWDG</b>	Tumor (NOS) - Custom Panel - 5 Probes	Johns Hopkins All Children's Hospital

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

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>87BHG</b>	OmniSeq INSIGHT	LabCorp
<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>7SS9G</b>	Lumera NGS Profile	Fulgent Genetics
<b>7SSAG</b>	Lumera Xpanded Profile	Fulgent Genetics
<b>6Y7FG</b>	OmniSeq Advance Assay	Labcorp   Oncology
<b>7VVDG</b>	HopeSeq Solid Tumors Comprehensive Panel	City of Hope National Medical Center - Molecular Diagnostic Laboratory
<b>7YCVG</b>	MI Cancer Seek + IHCs and Other Tests by Tumor Type	Caris Life Sciences

### Uveal Melanoma Prognostic Algorithmic Tests

<b>GTU</b>	<b>Test Name</b>	<b>Laboratory Name</b>
<b>6UTUG</b>	DecisionDx-UM	Castle Biosciences Inc
<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>7V35G</b>	Genomic Unity X-linked Intellectual Disability Plus Analysis	Variantyx, Inc.
<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>2BYZG</b>	X-linked Intellectual Disability (NGS Panel and Copy Number Analysis + Fragile X Repeat Expansion & Methylation)	MNG Laboratories
<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
<b>23XZG</b>	XLID (X-Linked Intellectual Disability) NGS Panel (Deletion/Duplication Only)	Fulgent Genetics
<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.

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