



## Genetic Tests and Laboratory Matrix

The matrix below contains all the genetic tests which Evolent (formerly National Imaging Associates, Inc.) manages on behalf of 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 [Texas Medicaid fee schedules](#) to determine which CPT codes are allowable for each genetic test managed by Evolent.**

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## Genetic Tests and Laboratory Matrix

### Arrhythmia Panel Tests

| GTU          | Test Name   | Laboratory Name  |
|--------------|---|--|
| <b>7XHCG</b> | Arrhythmia Comprehensive Panel                                    | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                  |
| <b>7VA6G</b> | Arrhythmia NGS Panel  | AiLife Diagnostics   |
| <b>7VA7G</b> | Arrhythmia NGS Panel Rapid  | AiLife Diagnostics   |
| <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>782SG</b> | Atrial fibrillation NGS Panel                                     | Connective Tissue Gene Tests   |
| <b>4JR5G</b> | Atrial Fibrillation NGS Panel (Deletion/Duplication Only)         | Fulgent Genetics   |
| <b>77VHG</b> | Atrial Fibrillation NGS Panel (Sequencing & Deletion/Duplication) | Fulgent Genetics   |
| <b>76WLG</b> | Atrial Fibrillation NGS Panel (Sequencing Only)                   | Fulgent Genetics   |
| <b>782TG</b> | Atrial fibrillation NGS Panel Next Generation Sequencing          | Connective Tissue Gene Tests   |
| <b>6MEFG</b> | Atrioventricular block NGS Panel                                  | Connective Tissue Gene Tests   |
| <b>6KDGG</b> | Atrioventricular block NGS Panel Deletion / Duplication           | Connective Tissue Gene Tests   |
| <b>6ME7G</b> | Atrioventricular block NGS Panel Next Generation Sequencing       | Connective Tissue Gene Tests   |
| <b>5TZFG</b> | Cardiac Arrhythmia Panel  | Northwest Clinical Genomics Lab  |
| <b>6MEDG</b> | Cardiac channelopathy NGS Panel                                   | Connective Tissue Gene Tests   |
| <b>6MALG</b> | Cardiac channelopathy NGS Panel Deletion / Duplication            | Connective Tissue Gene Tests   |
| <b>6MECG</b> | Cardiac channelopathy NGS Panel Next Generation Sequencing        | Connective Tissue Gene Tests   |
| <b>53VJG</b> | Comp Arrhythmia Panel   | Johns Hopkins Medical Institutions - Pathology Laboratory  |
| <b>7SCDG</b> | Comprehensive Arrhythmia Gene Panel, Varies                       | Mayo Clinic Laboratories   |

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

| GTU          | Test Name  | Laboratory Name  |
|--------------|--|--|
| <b>3E7RG</b> | Pan Arrhythmia (54 genes) Deletion/Duplication Panel | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |
| <b>2FQGG</b> | Pan Arrhythmia (54 genes) NGS + Sanger fill-in Panel | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |
| <b>54N6G</b> | Pan Arrhythmia (54 genes) NGS Panel                  | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |
| <b>7UY5G</b> | Postmortem Arrhythmia Gene Panel, Tissue             | Mayo Clinic Laboratories   |
| <b>2Z2MG</b> | RhythmNext   | Ambry Genetics   |
| <b>58GWG</b> | SCA Arrhythmia Panel                                 | Johns Hopkins Medical Institutions - Pathology Laboratory                    |
| <b>7B7BG</b> | STAT Comprehensive Arrhythmia Panel                  | PerkinElmer Genomics   |
| <b>6DTGG</b> | Sudden Cardiac Arrest Arrhythmia Panel               | GeneDx   |
| <b>77GKG</b> | Sudden Cardiac Arrest Panel                          | PreventionGenetics, part of Exact Sciences                                   |

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

| GTU          | Test Name  | Laboratory Name  |
|--------------|--|--|
| <b>2FHGG</b> | Autism NGS Panel (Deletion/Duplication Only)                                   | Fulgent Genetics   |
| <b>76VUG</b> | Autism NGS Panel (Sequencing & Deletion/Duplication)                           | Fulgent Genetics   |
| <b>78AYG</b> | Autism NGS Panel (Sequencing Only)   | Fulgent Genetics   |
| <b>78FSG</b> | Autism Spectrum Disorders (ASD) Panel  | PreventionGenetics, part of Exact Sciences   |
| <b>7BM2G</b> | AUTISM, INTELLECTUAL DISABILITY, and DEVELOPMENTAL DELAY GENE SEQUENCING PANEL | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| <b>76QWG</b> | AUTISM/AUTISM SPECTRUM DISORDER (53 gene panel)                                | Center for Human Genetics, Inc.  |
| <b>77KNG</b> | Autism/ID Panel  | GeneDx   |
| <b>78K3G</b> | Autism/ID Xpanded Panel  | GeneDx   |
| <b>77YKG</b> | AutismNext <sup>®</sup>  | Ambry Genetics   |
| <b>78FHG</b> | Autosomal Recessive Non-Specific Intellectual Disability Panel                 | The University of Chicago Genetic Services   |
| <b>7RW9G</b> | CHILD Neurodevelopmental Panel   | Children's Hospital Colorado Precision Diagnostics Laboratory                                    |

| GTU          | Test Name  | Laboratory Name                            |
|--------------|--|--|
| <b>77SAG</b> | Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA + Fragile X Repeat Expansion and Methylation) | MNG Laboratories                           |
| <b>77AUG</b> | Comprehensive Intellectual Disability / Autism (NGS Panel and Copy Number Analysis + mtDNA)  | MNG Laboratories                           |
| <b>787KG</b> | Focused Autism and Intellectual Disability Panel   | PerkinElmer Genomics                       |
| <b>76YDG</b> | Inherited Glycosylphosphatidylinositol Biosynthesis Defects (IGDs) Panel   | PreventionGenetics, part of Exact Sciences |
| <b>5FW2G</b> | Intellectual Disability Exome  | The University of Chicago Genetic Services |
| <b>5DPWG</b> | Intellectual Disability NGS Panel (Deletion/Duplication Only)  | Fulgent Genetics                           |
| <b>78F2G</b> | Intellectual Disability NGS Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics                           |
| <b>77GSG</b> | Intellectual Disability NGS Panel (Sequencing Only)  | Fulgent Genetics                           |
| <b>79Q6G</b> | Intellectual disability panel - NGS Panel (CNV included)   | Centogene                                  |
| <b>775YG</b> | Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Duo (IDEA panel of patient + 1 additional family members)          | PreventionGenetics, part of Exact Sciences |
| <b>783YG</b> | Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Family - Trio (IDEA panel of patient + 2 additional family members)         | PreventionGenetics, part of Exact Sciences |
| <b>77KFG</b> | Intellectual Disability, Epilepsy, and Autism (IDEA) Panel - Patient Only  | PreventionGenetics, part of Exact Sciences |
| <b>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> | PGmax™ - 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>7B62G</b> | STAT Focused Autism and Intellectual Disability Panel  | PerkinElmer Genomics                       |

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 77EPG | Syndromic Autism NGS Panel   | Greenwood Genetic Center - Molecular Diagnostic Laboratory |
| 77SFG | Syndromic Intellectual Disability (NGS Panel and Copy Number Analysis) | MNG Laboratories   |
| 78FXG | Top 99 Genetic Causes of Developmental Delay Panel                     | PreventionGenetics, part of Exact Sciences                 |

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

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

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

| GTU   | Test Name     | Laboratory Name          |
|-------|---------------|--------------------------|
| 74CFG | AlloMap Heart | Allina Health Laboratory |
| 525DG | AlloMap Heart | CareDx, Inc.             |

### BRCA1/2 Sequencing & Deletion/Duplication Tests

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 7SYQG | BRAC ANALYSIS,COMPREHENSIVE                                   | University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory |
| 6RPGG | BRACAnalysis CDx  | Myriad Genetics   |
| 2FFUG | BRCA AVANTAGE,COMPREHENSIVE                                   | Empire City Laboratories, Inc.  |
| 3GPZG | BRCA Comprehensive (BRCA 1 and BRCA 2 Seq & Del/Dup Analysis) | Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories                         |
| 26REG | BRCA Panel (BRCA1, BRCA2)                                     | Palo Verde Laboratory - division of Sonora Quest Lab  |
| 2E63G | BRCA Panel (BRCA1, BRCA2)                                     | Quest Diagnostics   |
| 7V2JG | BRCA1 & BRCA2 Panel   | Baylor Genetics, LLC  |



| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 55MGG | BRCA1 AND BRCA2 ANALYSIS (SEQUENCING & MLPA FOR BOTH GENES)                                | Center for Human Genetics, Inc.   |
| 5BRJG | BRCA1 and BRCA2 Focus (Sequencing & Deletion/Duplication)                                  | Fulgent Genetics  |
| 6U9AG | BRCA1 and BRCA2 Focus Panel (Germline)   | NeoGenomics Laboratories  |
| 7U27G | BRCA1 and BRCA2 Sequencing and Del/Dup (NGS)   | University of Michigan - Michigan Medical Genetics Laboratories                   |
| 2ELCG | BRCA1 and BRCA2 Sequencing and Deletion/Duplication  | Knight Molecular Diagnostic   |
| 7QGJG | BRCA1 AND BRCA2, COMPREHENSIVE   | Clinical Pathology Laboratories   |
| 6UQAG | BRCA1 and BRCA2-Associated HBOC Syndrome Panel, Sequencing and Deletion/Duplication        | ARUP Laboratories   |
| 6LTEG | BRCA1&2 Analysis   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 79AHG | BRCA1, BRCA2 panel Combi (with MLPA) - NGS Panel (MLPA included)                           | Centogene   |
| 79AJG | BRCA1, BRCA2 panel Plus - NGS Panel (CNV included)   | Centogene   |
| 2YYZG | BRCA1/2 seq and del/dup  | Ambry Genetics  |
| 4RLXG | BRCA1/2 Sequencing and Del/Dup Analysis  | BioReference Laboratories   |
| 6DTQG | BRCA1/2 Sequencing and Del/Dup Analysis  | GeneDx  |
| 3D8PG | BRCA1/2 Sequencing and Deletion Duplication Analysis                                       | ACL Laboratories  |
| 5FUUG | BRCA1/2: Comprehensive BRCA Analysis by Gene Sequencing with Deletion/Duplication Analysis | Medical Diagnostic Laboratories, LLC  |
| 7DLBG | BRCA1/BRCA2 Genes, Full Gene Analysis, Varies  | Mayo Clinic Laboratories  |
| 6LS6G | BRCAssure : BRCA1 and BRCA2 Comprehensive Analysis   | LabCorp   |
| 546VG | BRCAssure: BRCA1 / 2 Comprehensive Analysis  | Labcorp   Oncology  |
| 58G6G | BRCAssure: Comprehensive BRCA1 / 2 Analysis  | Integrated Genetics   |
| 2C88G | Empower - BRCA1 & BRCA2 Hereditary Cancer Test   | Natera  |
| 7X27G | GxVISION Hereditary Cancer Risk Assessment BRCA1/2 Genes                                   | Otogenetics   |
| 57DKG | Hereditary Breast and Ovarian Cancer BRCA1/2 Panel   | PreventionGenetics, part of Exact Sciences  |
| 32SZG | Hereditary Breast and Ovarian Cancer Syndrome Panel  | PerkinElmer Genomics  |
| 7WXHG | Hereditary cancer BRCA1 BRCA2  | University of Minnesota Physicians Outreach Laboratory                            |

| GTU   | Test Name  | Laboratory Name                     |
|-------|--|-------------------------------------|
| 2V8RG | Integrated BRAC Analysis, EDTA Whole Blood (BRACA 1 & 2 Testing) | Marshfield Labs                     |
| 6RMHG | Integrated BRACAnalysis  | Myriad Genetics                     |
| 7QDVG | Integrated Reflex BRACAnalysis                                   | Myriad Genetics                     |
| 442GG | Invitae BRCA1 and BRCA2 Panel                                    | Invitae Corporation                 |
| 44N2G | Invitae BRCA1 and BRCA2 STAT Panel                               | Invitae Corporation                 |
| 7V3GG | OnkoRisk BRCA Panel (Non-NYS)                                    | BioReference Laboratories           |
| 7SLMG | PrevenTest, Custom (BRCA1, BRCA2)                                | Advanced Molecular Diagnostics, LLC |
| 6RN9G | Reflex BRACAnalysis  | Myriad Genetics                     |
| 7B7QG | STAT Hereditary Breast and Ovarian Cancer Syndrome Panel         | PerkinElmer Genomics                |

### BRCA1/2 Sequencing Tests

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

### Breast Cancer Prognostic Algorithmic Tests

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

| GTU   | Test Name  | Laboratory Name |
|-------|--|-----------------|
| 2KTBG | Prosigna Breast Cancer Prognostic Gene Signature Assay | Veracyte        |

### Breast Cancer Treatment and Prognostic Algorithmic Tests

| GTU   | Test Name                                | Laboratory Name                  |
|-------|--|----------------------------------|
| 2VDEG | Oncotype Dx Breast Cancer Assay          | Marshfield Labs                  |
| 7BNGG | Oncotype DX Breast Recurrence Score Test | Exact Sciences Laboratories, LLC |

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

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

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

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

| GTU   | Test Name    | Laboratory Name |
|-------|--------------|-----------------|
| 7UQXG | Plasma Focus | LabCorp         |

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

| GTU   | Test Name                              | Laboratory Name      |
|-------|--|----------------------|
| 2AHTG | OncoBEAM™ CRC1: KRAS, NRAS, BRAF, HRAS | Sysmex Inostics, INC |

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

| GTU   | Test Name                                  | Laboratory Name          |
|-------|--|--------------------------|
| 7SGYG | CNSide Biomarker Profile, Lung             | Biocept                  |
| 7ACFG | GeneStrat                                  | Biodesix                 |
| 7ACGG | GeneStrat NGS                              | Biodesix                 |
| 4764G | Genestrat Test                             | ACL Laboratories         |
| 7TRFG | InVisionFirst -Lung Liquid Biopsy          | Inivata Ltd              |
| 4FGPG | InVisionFirst -Lung Liquid Biopsy          | NeoGenomics Laboratories |
| 7RVXG | IQlung Treatment Guidance Testing          | Biodesix                 |
| 28XAG | OncoBEAM™ Lung2: EGFR, KRAS, BRAF          | Sysmex Inostics, INC     |
| 6UU4G | Resolution ctDx Lung assay (Liquid Biopsy) | Labcorp   Oncology       |

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

| GTU   | Test Name  | Laboratory Name                                       |
|-------|--|---|
| 2KSUG | 511810 Follow-Up Prenatal CMA qPCR   | LabCorp   |
| 79AMG | CentoArrayCyto 750K - Array CGH (prenatal)   | Centogene   |
| 79APG | CentoArrayCyto HD - Array CGH (prenatal)   | Centogene   |
| 2EKCG | Chromosomal Microarray - Prenatal Diagnosis  | Knight Molecular Diagnostic                           |
| 2LBVG | Chromosomal Microarray Analysis - Prenatal CMA Amniotic Fluid (Affymetrix CytoScan HD array)           | Washington University in St. Louis Pathology Services |
| 2QYYG | Chromosomal Microarray Analysis - Prenatal CMA Chorionic Villi Sampling (Affymetrix CytoScan HD array) | Washington University in St. Louis Pathology Services |
| 33X4G | Chromosomal Microarray Prenatal (CMAP)   | Marshfield Labs                                       |

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 2ETFG | Chromosomal Microarray, Prenatal, ClariSure                                | Johns Hopkins Medical Institutions - Pathology Laboratory                                    |
| 6LEBG | Chromosomal Microarray, Prenatal   | Hennepin County Medical Center   |
| 6R47G | Chromosomal Microarray, Prenatal, Amniotic Fluid/Chorionic Villus Sampling | Mayo Clinic Laboratories   |
| 7U36G | Chromosomal Microarray, Prenatal, Amniotic Fluid/Chorionic Villus Sampling | University of Michigan - Michigan Medical Genetics Laboratories                              |
| 2EQ3G | Chromosomal Microarray, Prenatal, ClariSure Oligo-SNP                      | Quest Diagnostics  |
| 2WTCG | CHROMOSOMAL MICROARRAY,PRENATAL CLARISURE                                  | Empire City Laboratories, Inc.   |
| 7U5HG | Cytogenomic SNP Microarray&nbsp;-&nbsp;Fetal                               | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory               |
| 25DWG | Cytogenomic SNP Microarray, Fetal  | ARUP Laboratories  |
| 4XSNG | Cytogenomic SNP Microarray, Fetal  | TriCore Reference Laboratories   |
| 3DQXG | Cytogenomic SNP Microarray, Fetal  | UCSD - Molecular Diagnostics & Cytogenetics Laboratory                                       |
| 78KYG | Cytogenomic SNP Microarray, Fetal  | University of California Davis Health System - UCDCMC - Molecular and Cytogenetic Laboratory |
| 2P6UG | Expanded Chromosomal Microarray Analysis - Prenatal - Amniotic Fluid       | Baylor Genetics, LLC   |
| 4JAVG | Expanded Chromosomal Microarray Analysis - Prenatal - CVS                  | Baylor Genetics, LLC   |
| 5BR7G | Genomic Microarray, Prenatal (Amniotic Fluid or CVS)                       | ACL Laboratories   |
| 7V33G | IriSight™? CNV AnalysisAccepted Specimens                                  | Variyantx, Inc.  |
| 7X43G | POC Microarray with 5-Cell Chromosome Analysis                             | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory              |
| 328BG | Prenatal Cytogenomic Microarray  | University Hospitals   |
| 2LEXG | Prenatal Microarray  | Greenwood Genetic Center - Molecular Diagnostic Laboratory                                   |
| 7X46G | Prenatal Microarray with 5-Cell Chromosome Analysis                        | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory              |
| 7X49G | Prenatal Microarray with Parental Testing                                  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory              |
| 7X45G | Prenatal Microarray without Parental Testing                               | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory              |
| 7C4VG | Prenatal Whole Genome Chromosomal Microarray                               | BioReference Laboratories  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 6WNXG | Prenatal Whole Genome Chromosomal Microarray                            | GeneDx  |
| 6LUQG | Rapid Prenatal Chromosomal Microarray via aCGH and SNP - Prenatal Test  | PreventionGenetics, part of Exact Sciences                                      |
| 7C4XG | Reflex to Prenatal Whole Genome Chromosomal Microarray from Chromosomes | BioReference Laboratories   |
| 5BNSG | Reveal SNP Microarray – Prenatal  | Integrated Genetics   |
| 2EPWG | SNP Array for Prenatal Analysis (aka Microarray)                        | UCSF Molecular Diagnostics Laboratory   |
| 7X47G | SNP Microarray  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 3U6CG | SNP Microarray (Direct)-Prenatal (Reveal)                               | LabCorp   |
| 3N8VG | SNP Microarray-Prenatal (Reveal)  | LabCorp   |

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

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 6S7ZG | Anora Miscarriage Test (Fresh)  | Natera  |
| 2EXPG | Anora Miscarriage Test (Paraffin)   | Natera  |
| 2EW5G | CGH-POC: Comparative Genomic Hybridization  | New Jersey Medical School - Institute of Genomic Medicine       |
| 2EDZG | Chromosomal Microarray Analysis - CMA Products of Conception (POC) (Affymetrix CytoScan HD array) | Washington University in St. Louis Pathology Services           |
| 6LRAG | Chromosomal Microarray Analysis (CMA) - Products of Conception                                    | Allina Health Laboratory  |
| 6R2VG | Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth                            | Mayo Clinic Laboratories  |
| 7U37G | Chromosomal Microarray, Autopsy, Products of Conception, or Stillbirth                            | University of Michigan - Michigan Medical Genetics Laboratories |
| 6R3NG | Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue                         | Mayo Clinic Laboratories  |
| 7U35G | Chromosomal Microarray, Autopsy/Products of Conception/Stillbirth, Tissue                         | University of Michigan - Michigan Medical Genetics Laboratories |
| 2E8MG | Chromosomal Microarray, POC, ClariSure Oligo-SNP  | Quest Diagnostics   |
| 3PJJG | CHROMOSOMAL MICROARRAY, POC, CLARISURE OLIGO-SNP, POC   | Empire City Laboratories, Inc.                                  |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 3KVVG | Comparative Genomic Hybridization (CGH): Products of Conception (POC)            | New Jersey Medical School - Institute of Genomic Medicine                       |
| 7DK8G | Cytogenomic Molecular Inversion Probe Array FFPE Tissue - Products of Conception | ARUP Laboratories   |
| 2EKBG | Genomic Microarray, POC  | ACL Laboratories  |
| 2EXDG | Genomic SNP Microarray, Products of Conception                                   | ARUP Laboratories   |
| 7PTBG | Microarray Analysis - PREGNANCY LOSS   | Nebraska Medical Center - Molecular Diagnostic Laboratory                       |
| 2ENDG | Microarray-Products of Conception (POC) Reveal FFPE                              | LabCorp   |
| 2E65G | Microarray-Products of Conception (POC) Reveal FFPE, Data Transfer               | LabCorp   |
| 7XJKG | POC (Products of Conception) Microarray Analysis                                 | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 2M7XG | Pregnancy Loss Chromosomal SNP Microarray  | Colorado Genetics Laboratory  |
| 3JXFG | Products of Conception Microarray + SNP  | Cleveland Clinic Laboratories   |
| 4ZPRG | Reveal SNP Microarray – POC  | Integrated Genetics   |
| 5FU7G | SNP Array for Tissue and POC (aka Microarray)                                    | UCSF Molecular Diagnostics Laboratory   |
| 34WFG | SNP Microarray – Products of Conception  | Knight Molecular Diagnostic   |
| 3U6BG | SNP Microarray Products of Conception (POC) / Tissue (Reveal )                   | LabCorp   |

### Chromosomal Microarray (SNP and CGH) Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 25SKG | 2.7 M SNP Chromosomal Microarray                          | Nicklaus Children's Hospital - Miami Genetic Laboratories            |
| 4LCPG | Array CGH   | UAB Cytogenetics Laboratory  |
| 3R4MG | Array Comparative Genomic Hybridization                   | Children's Hospital and Research Center Oakland - Molecular Genetics |
| 2KYRG | Array-based Comparative Genome Hybridization, Genetic Dx. | Stanford Clinical Laboratories - Biochemical Genetics Laboratory     |
| 3GPJG | Assure SNP Microarray Analysis (FDA cleared)              | CytoGenX   |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 2E8LG | Assure SNP Microarray Analysis (FDA cleared) – Prenatal & Postnatal                   | CytoGenX  |
| 7XJLG | Autopsy Microarray Analysis   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory           |
| 3AG7G | CGH: Comparative Genomic Hybridization  | New Jersey Medical School - Institute of Genomic Medicine                                 |
| 3F5GG | CHROMOSOMAL MICROARRAY  | Detroit Medical Center University Laboratories - Molecular Genetics Diagnostic Laboratory |
| 4CURG | CHROMOSOMAL MICROARRAY  | Fullerton Genetics Center - Mission Health  |
| 5JPRG | Chromosomal Microarray - Postnatal  | Knight Molecular Diagnostic   |
| 6XNCG | Chromosomal Microarray (MicroarrayDx)   | GeneDx  |
| 687XG | Chromosomal Microarray Analysis   | Center for Genetic Testing at Saint Francis   |
| 38VKG | Chromosomal Microarray Analysis   | UCLA Diagnostic Molecular Pathology Laboratory  |
| 3UXBG | Chromosomal Microarray Analysis - CMA Fibroblasts (Affymetrix CytoScan HD array)      | Washington University in St. Louis Pathology Services                                     |
| 3HN3G | Chromosomal Microarray Analysis - CMA Peripheral Blood (Affymetrix CytoScan HD array) | Washington University in St. Louis Pathology Services                                     |
| 6NL2G | Chromosomal Microarray Analysis - HR  | Baylor Genetics, LLC  |
| 2ZAVG | Chromosomal Microarray Analysis - HR + SNP Screen (Comprehensive)                     | Baylor Genetics, LLC  |
| 6LRBG | Chromosomal microarray analysis (CMA) - Blood   | Allina Health Laboratory  |
| 4DNFG | Chromosomal Microarray Analysis for Constitutional Abnormalities                      | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory         |
| 2EW7G | Chromosomal Microarray only   | Washington University in St. Louis Pathology Services                                     |
| 53JEG | Chromosomal Microarray SNP, Constitutional  | Cleveland Clinic Laboratories   |
| 3A53G | Chromosomal Microarray with abbreviated karyotype                                     | Washington University in St. Louis Pathology Services                                     |
| 32EDG | Chromosomal Microarray with concurrent karyotype                                      | Washington University in St. Louis Pathology Services                                     |
| 333GG | Chromosomal Microarray with reflex karyotype  | Washington University in St. Louis Pathology Services                                     |
| 4XSMG | Chromosomal Microarray, Congenital, Bld (CMACB)                                       | Marshfield Labs   |
| 6R2TG | Chromosomal Microarray, Congenital, Blood   | Mayo Clinic Laboratories  |
| 7R9LG | Chromosomal Microarray, Hematologic Malignancy, ClariSure Oligo-SNP (90961)           | Rady Children's Institute for Genomic Medicine  |



| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 2Z55G | Chromosomal Microarray, POC FFPE, ClariSure Oligo-SNP                           | Quest Diagnostics   |
| 8EJVG | Chromosomal Microarray, POC, ClariSure Oligo-SNP                                | Medfusion   |
| 8EPBG | Chromosomal Microarray, Postnatal Familial Follow-up, ClariSure Oligo-SNP       | Quest Diagnostics   |
| 2ETDG | Chromosomal Microarray, Postnatal, ClariSure Oligo-SNP                          | Quest Diagnostics   |
| 2E8NG | Chromosomal Microarray, Postnatal, ClariSure Oligo-SNP (16478)                  | Rady Children's Institute for Genomic Medicine  |
| 2EP9G | Chromosomal SNP Microarray  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics   |
| 2BG9G | Chromosomal SNP Microarray  | Seattle Children's Hospital   |
| 68Y6G | Chromosome Array (aCGH)   | Center for Genetic Testing at Saint Francis   |
| 2DXNG | Chromosome Microarray   | Palo Verde Laboratory - division of Sonora Quest Lab  |
| 53TQG | Chromosome Microarray   | University of Virginia Health System  |
| 54CPG | Chromosome Microarray Analysis  | Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital                             |
| 2ELZG | Chromosome Microarray Analysis - Constitutional                                 | John Hopkins All Children's Hospital  |
| 7S97G | Chromosome Microarray, Congenital   | Sanford USD Medical Center - Sanford Clinic USD Genetics Laboratory   |
| 2DYKG | CHROMOSOME MICROARRAY, POSTNATAL  | Empire City Laboratories, Inc.  |
| 7QHSG | CHROMOSOME SNP MICROARRAY   | University of Texas Medical Branch - UTMB - Porphyria Laboratory  |
| 5FQBG | Chromosome Specific Interphase FISH   | Center for Genetic Testing at Saint Francis   |
| 7X4CG | CNS Tumor Classification by Methylation Array                                   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                                     |
| 2LLCG | Combimatrix Combi SNP Array Tissue  | Norton CPA Lab  |
| 2FNJG | Compar Genomic Hybrid-SNP   | Norton CPA Lab  |
| 2EPKG | Comparative Genomic Hybridization (CGH):Mental Retardation/ Developmental Delay | New Jersey Medical School - Institute of Genomic Medicine   |
| 7WJHG | Constitutional Chromosomal Microarray (Copy Number)                             | University of Minnesota Physicians Outreach Laboratory  |
| 6T6QG | Constitutional Chromosomal Microarray Analysis – Peripheral Blood               | Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine |
| 2E7LG | Constitutional Chromosomal Microarray Analysis (CMA)                            | Indiana University School of Medicine - Cytogenetics Laboratory   |
| 7WBMG | Constitutional Limited Chromosomal Microarray (Copy number only) (Charged)      | University of Minnesota Physicians Outreach Laboratory  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 7WPQG | Constitutional or Products of Conception (POC) Chromosomal Microarray (Copy Number/SNP)   | University of Minnesota Physicians Outreach Laboratory                                      |
| 2ETHG | Constitutional SNP Array Karyotyping, Constitutional Chromosome Microarray Analysis (CMA) | Beaumont Laboratories - Molecular Pathology Lab   |
| 2M4WG | Cytogenomic Microarray Analysis   | University of Washington Medicine - Pathology - Cytogenetics and Genomics Laboratory        |
| 49JCG | Cytogenomic Microarray Analysis of Postnatal Blood  | Akron Children's Hospital   |
| 7W9QG | Cytogenomic Microarray SNP Fetal  | University of Minnesota Physicians Outreach Laboratory                                      |
| 49W6G | Cytogenomic SNP array (postnatal)   | The University of Chicago Genetic Services  |
| 2E3BG | Cytogenomic SNP Microarray  | ARUP Laboratories   |
| 8EBHG | Cytogenomic SNP Microarray  | Medfusion   |
| 2MTFG | Cytogenomic SNP Microarray  | UCSD - Molecular Diagnostics & Cytogenetics Laboratory                                      |
| 2X3MG | Cytogenomic SNP Microarray  | University of California Davis Health System - UCDCM - Molecular and Cytogenetic Laboratory |
| 7XL3G | Cytogenomic SNP Microarray  | University of Illinois at Chicago - Biochemical Genetics Laboratory                         |
| 7U9JG | Cytogenomic SNP Microarray  | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory              |
| 2542G | Cytogenomic SNP Microarray Buccal Swab  | ARUP Laboratories   |
| 2D5QG | FirstStepDx PLUS  | Bionano Laboratories  |
| 7UHHG | Genomic Microarray Analysis - Parental study, Genome Wide Array                           | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory              |
| 3MA8G | Genomic Microarray, Blood   | ACL Laboratories  |
| 2ENHG | Genomic SNP Microarray, Products of Conception  | University of California Davis Health System - UCDCM - Molecular and Cytogenetic Laboratory |
| 7UHKG | Genomic SNP Microarray, Products of Conception  | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory              |
| 2ETGG | Illumina Global Screening Array Sequencing Test   | Gene by Gene  |
| 2EQNG | Illumina Microarray Analysis  | University of Wisconsin - Madison WSLH - UW Cytogenetic Services                            |
| 2EMZG | Illumina Microarray Analysis- Targeted Family   | University of Wisconsin - Madison WSLH - UW Cytogenetic Services                            |
| 7TG6G | Invitae Chromosomal Microarray Analysis (CMA)   | Invitae Corporation   |

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

| GTU   | Test Name  | Laboratory Name                                     |
|-------|--|---|
| 2Z5JG | SNP Microarray Pediatric   | Integrated Regional Laboratories                    |
| 2EKEG | SNP microarray Processing, Extraction and Storage                                    | UCSF Molecular Diagnostics Laboratory               |
| 49AAG | SNP Microarray when Routine Chrom or High Res and Fragile X have ALREADY been billed | Center for Human Genetics, Inc.                     |
| 2EQQG | SNP Microarray-Pediatric (Reveal)  | LabCorp   |
| 52MEG | SNP Oligonucleotide Microarray Analysis (SOMA)                                       | Columbia University - Personalized Genomic Medicine |
| 6NNKG | Whole-Genome Chromosomal Microarray (CMA-ISCA) via the aCGH and SNP Test #2000       | PreventionGenetics, part of Exact Sciences          |

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

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

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

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 23DQG | Cardiomyopathy and Arrhythmia Panel, Sequencing and Deletion/Duplication  | ARUP Laboratories                                      |
| 2DTLG | Cardiomyopathy and Arrhythmia Sequencing Panel  | UCSD - Molecular Diagnostics & Cytogenetics Laboratory |
| 7TP5G | CardioNext: Arrhythmia  | Ambry Genetics   |
| 77BXG | CardioNext: Cardiomyopathy  | Ambry Genetics   |
| 2LRSG | Combined Cardiac Panel  | GeneDx   |
| 7SCFG | Comprehensive Arrhythmia and Cardiomyopathy Gene Panel, Varies  | Mayo Clinic Laboratories                               |
| 3FQNG | Comprehensive Cardiac Arrhythmia/Cardiomyopathy Panel   | Northwest Clinical Genomics Lab                        |
| 438XG | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel   | Invitae Corporation                                    |
| 7DR2G | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel - UNLOCKCARDIO  | Invitae Corporation                                    |
| 43H3G | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy | Invitae Corporation                                    |

| GTU   | Test Name  | Laboratory Name          |
|-------|--|--------------------------|
| 7QKVG | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Arrhythmia and Cardiomyopathy - UNLOCKCARDIO | Invitae Corporation      |
| 43FSG | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes                           | Invitae Corporation      |
| 7QL3G | Invitae Arrhythmia and Cardiomyopathy Comprehensive Panel-Add-on Sudden Unexpected Death in Epilepsy (SUDEP) Genes - UNLOCKCARDIO            | Invitae Corporation      |
| 7UY6G | Postmortem Cardiomyopathy and Arrhythmia Gene Panel, Tissue  | Mayo Clinic Laboratories |
| 4XFMG | Sudden Death Syndrome NGS Panel (Deletion/Duplication Only)  | Fulgent Genetics         |
| 77CZG | Sudden Death Syndrome NGS Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics         |
| 77U8G | Sudden Death Syndrome NGS Panel (Sequencing Only)  | Fulgent Genetics         |

### Comprehensive Cardiomyopathy Panel Tests

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 7VAEG | Cardiomyopathy NGS Panel   | AiLife Diagnostics  |
| 7VAFG | Cardiomyopathy NGS Panel Rapid   | AiLife Diagnostics  |
| 4MUVG | Cardiomyopathy Panel   | Northwest Clinical Genomics Lab                                       |
| 7DMVG | Cardiomyopathy Panel   | Washington University in St. Louis Genomics and Pathology Services    |
| 45LWG | Cardiomyopathy Pediatric Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 79PLG | CentoCardio - NGS Panel (CNV included)                                     | Centogene   |
| 2YZFG | CMNext   | Ambry Genetics  |
| 28S8G | Comp Cardiomyopathy Panel  | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 2BXWG | Comprehensive Cardiomyopathy (NGS Panel and Copy Number Analysis + mtDNA)  | MNG Laboratories  |
| 7SCCG | Comprehensive Cardiomyopathy Gene Panel, Varies                            | Mayo Clinic Laboratories  |
| 3XZHG | Comprehensive Cardiomyopathy NGS Panel (Deletion/Duplication Only)         | Fulgent Genetics  |
| 25DLG | Comprehensive Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication) | Fulgent Genetics  |

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 33E5G | Comprehensive Cardiomyopathy NGS Panel (Sequencing Only)  | Fulgent Genetics   |
| 345YG | Comprehensive Cardiomyopathy Panel  | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 332EG | Comprehensive Cardiomyopathy Panel  | Knight Molecular Diagnostic  |
| 7DQHG | Comprehensive Cardiomyopathy Panel  | PerkinElmer Genomics   |
| 56ZPG | GeneSeq Cardio Familial Cardiomyopathy Panel  | LabCorp  |
| 3GBZG | GeneSeq : Cardio-Familial Cardiomyopathy Profile  | Integrated Genetics  |
| 5FWFG | Invitae Cardiomyopathy Comprehensive Panel  | Invitae Corporation  |
| 3YNXG | Invitae Cardiomyopathy Comprehensive Panel - LYSO   | Invitae Corporation  |
| 2C5NG | Invitae Cardiomyopathy Comprehensive Panel - TTR  | Invitae Corporation  |
| 7D5WG | Invitae Cardiomyopathy Comprehensive Panel - TTRCA  | Invitae Corporation  |
| 24WTG | Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy         | Invitae Corporation  |
| 3YRRG | Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy - LYSO  | Invitae Corporation  |
| 2C7JG | Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy - TTR   | Invitae Corporation  |
| 7QLCG | Invitae Cardiomyopathy Comprehensive Panel-Add-on Preliminary-evidence Genes for Cardiomyopathy - TTRCA | Invitae Corporation  |
| 5U3FG | Pan Cardiomyopathy (92 genes) Deletion/Duplication Panel  | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 2G3NG | Pan Cardiomyopathy (92 genes) NGS + Sanger fill-in Panel  | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 5638G | Pan Cardiomyopathy (92 genes) NGS Panel   | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 7BLQG | Pan Cardiomyopathy Panel  | Johns Hopkins Medical Institutions - Pathology Laboratory  |
| 57C9G | Pan Cardiomyopathy Panel  | PreventionGenetics, part of Exact Sciences   |
| 7UHG  | Pan Cardiomyopathy Panel (62 Genes)   | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory                   |
| 7UY7G | Postmortem Cardiomyopathy Gene Panel, Tissue  | Mayo Clinic Laboratories   |
| 7B77G | STAT Comprehensive Cardiomyopathy Panel   | PerkinElmer Genomics   |

| GTU   | Test Name                           | Laboratory Name |
|-------|-------------------------------------|-----------------|
| 7AP3G | Suggested Custom Slice - Lymphedema | GeneDx          |

### Dilated Cardiomyopathy (DCM) Panel Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 2YW9G | DCMNext   | Ambry Genetics   |
| 5NAGG | Dilated & Arrhythmogenic Cardiomyopathy NGS Panel   | Greenwood Genetic Center - Molecular Diagnostic Laboratory                                       |
| 3JZVG | Dilated Cardiomyopathy – DCM (52 genes) Deletion/Duplication Panel                          | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 2FP7G | Dilated Cardiomyopathy – DCM (52 genes) NGS + Sanger fill-in Panel                          | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 3CBJG | Dilated Cardiomyopathy – DCM (52 genes) NGS Panel   | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 3XP4G | Dilated Cardiomyopathy (DCM) Panel  | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 7DMXG | Dilated Cardiomyopathy (DCM) Panel  | Washington University in St. Louis Genomics and Pathology Services                               |
| 7SC4G | Dilated Cardiomyopathy and Left Ventricular Noncompaction Cardiomyopathy Gene Panel, Varies | Mayo Clinic Laboratories   |
| 44D6G | Dilated Cardiomyopathy NGS Panel (Deletion/Duplication Only)                                | Fulgent Genetics   |
| 25DFG | Dilated Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication)                        | Fulgent Genetics   |
| 3CJ3G | Dilated Cardiomyopathy NGS Panel (Sequencing Only)  | Fulgent Genetics   |
| 3CBUG | Dilated Cardiomyopathy Panel  | Knight Molecular Diagnostic  |
| 7XHEG | Dilated Cardiomyopathy Panel  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                  |
| 3XNVG | Dilated Cardiomyopathy Panel  | PerkinElmer Genomics   |
| 5CZ6G | Dilated Cardiomyopathy Panel  | PreventionGenetics, part of Exact Sciences   |
| 2KTAG | Dilated Cardiomyopathy Panel, Sequencing  | ARUP Laboratories  |
| 3XPBG | Dilated Cardiomyopathy Sequencing Panel with CNV Detection: Institution-Specific            | PreventionGenetics, part of Exact Sciences   |
| 2KYQG | Invitae Dilated Cardiomyopathy and Left Ventricular Noncompaction Panel                     | Invitae Corporation  |

| GTU          | Test Name   | Laboratory Name      |
|--------------|---|----------------------|
| <b>5FWEG</b> | Invitae Dilated Cardiomyopathy and Left Ventricular Noncompaction Panel-Add-on Preliminary-evidence Genes for Dilated Cardiomyopathy and Left Ventricular Noncompaction | Invitae Corporation  |
| <b>7B78G</b> | STAT Dilated Cardiomyopathy Panel   | PerkinElmer Genomics |

## Dystonia Panel Tests

| GTU          | Test Name   | Laboratory Name                            |
|--------------|---|--|
| <b>77W5G</b> | Basal Ganglia Calcification Dystonia (NGS Panel and Copy Number Analysis)                           | MNG Laboratories                           |
| <b>5X7HG</b> | Complete Dopa-Responsive Dystonia (DYT5) Evaluation   | Athena Diagnostics Inc                     |
| <b>78DLG</b> | Comprehensive Dystonia (NGS Panel and Copy Number Analysis + mtDNA + HTT Repeat Expansion Analysis) | MNG Laboratories                           |
| <b>77FMG</b> | Comprehensive Dystonia (NGS Panel and Copy Number Analysis + mtDNA)                                 | MNG Laboratories                           |
| <b>249UG</b> | Dopa-Responsive Dystonia (NGS Panel and Copy Number Analysis)                                       | MNG Laboratories                           |
| <b>6FW7G</b> | Dopa-Responsive Dystonia NGS Panel (Deletion/Duplication Only)                                      | Fulgent Genetics                           |
| <b>6CT3G</b> | Dopa-Responsive Dystonia NGS Panel (Sequencing & Deletion/Duplication)                              | Fulgent Genetics                           |
| <b>3PSFG</b> | Dopa-Responsive Dystonia NGS Panel (Sequencing Only)  | Fulgent Genetics                           |
| <b>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>775XG</b> | Dystonia Panel  | GeneDx                                     |
| <b>775BG</b> | Dystonia Panel  | Knight Molecular Diagnostic                |
| <b>77JMG</b> | Dystonia Panel  | PerkinElmer Genomics                       |
| <b>78FUG</b> | Dystonia Panel  | PreventionGenetics, part of Exact Sciences |
| <b>778BG</b> | Invitae Dystonia Comprehensive Panel  | Invitae Corporation                        |



| GTU   | Test Name   | Laboratory Name        |
|-------|---|------------------------|
| 76S3G | Invitae Dystonia Comprehensive Panel-Add-on Preliminary-evidence Genes for Dystonia | Invitae Corporation    |
| 4SRKG | Isolated Dystonia Evaluation  | Athena Diagnostics Inc |
| 5X6QG | Myoclonic Dystonia Panel  | PerkinElmer Genomics   |
| 77MVG | OXPHOS Defect Dystonia (NGS Panel and Copy Number Analysis + mtDNA)                 | MNG Laboratories       |
| 76Q3G | Primary Dystonia (NGS Panel and Copy Number Analysis)                               | MNG Laboratories       |
| 7AYEG | STAT Dystonia Panel   | PerkinElmer Genomics   |
| 7AYCG | STAT Myoclonic Dystonia Panel   | PerkinElmer Genomics   |

### Epilepsy and Seizure Disorder Panel Tests

| GTU   | Test Name   | Laboratory Name             |
|-------|---|-----------------------------|
| 65CUG | Actionable Epilepsy NGS Panel (Deletion/Duplication Only)               | Fulgent Genetics            |
| 78D5G | Actionable Epilepsy NGS Panel (Sequencing & Deletion/Duplication)       | Fulgent Genetics            |
| 76Y6G | Actionable Epilepsy NGS Panel (Sequencing Only)                         | Fulgent Genetics            |
| 3UT6G | Adolescent/Adult Epilepsy NGS Panel (Deletion/Duplication Only)         | Fulgent Genetics            |
| 78BJG | Adolescent/Adult Epilepsy NGS Panel (Sequencing & Deletion/Duplication) | Fulgent Genetics            |
| 77UVG | Adolescent/Adult Epilepsy NGS Panel (Sequencing Only)                   | Fulgent Genetics            |
| 3PZSG | Childhood Epilepsy NGS Panel (Deletion/Duplication Only)                | Fulgent Genetics            |
| 77UAG | Childhood Epilepsy NGS Panel (Sequencing & Deletion/Duplication)        | Fulgent Genetics            |
| 76XSG | Childhood Epilepsy NGS Panel (Sequencing Only)                          | Fulgent Genetics            |
| 784QG | Childhood Epilepsy Panel  | Knight Molecular Diagnostic |
| 87ASG | Clinical Epilepsy NGS Panel   | LabCorp                     |
| 78L9G | Clinical Epilepsy NGS Panel   | MNG Laboratories            |
| 77ASG | Comprehensive Epilepsy (NGS Panel and Copy Number Analysis + mtDNA)     | MNG Laboratories            |
| 73X4G | Comprehensive Epilepsy NGS Panel  | LabCorp                     |

| <b>GTU</b>   | <b>Test Name</b>  | <b>Laboratory Name</b>                                     |
|--------------|---|--|
| <b>78KDG</b> | Comprehensive Epilepsy Panel  | GeneDx   |
| <b>778XG</b> | Comprehensive Epilepsy Panel  | PerkinElmer Genomics                                       |
| <b>7UUBG</b> | Comprehensive Epilepsy Panel, Sequencing and Deletion/Duplication   | ARUP Laboratories  |
| <b>7BNUG</b> | Comprehensive Epilepsy With or Without Encephalopathy Gene Panel, Varies  | Mayo Clinic Laboratories                                   |
| <b>78FGG</b> | Early Infantile Epileptic Encephalopathy (EIEE) Panel   | The University of Chicago Genetic Services                 |
| <b>77V3G</b> | Early Infantile Epileptic Encephalopathy NGS Panel  | Greenwood Genetic Center - Molecular Diagnostic Laboratory |
| <b>77FPG</b> | Early Infantile Epileptic Encephalopathy Panel  | PreventionGenetics, part of Exact Sciences                 |
| <b>5DRKG</b> | Early Onset Epileptic Encephalopathy NGS Panel (Deletion/Duplication Only)                                      | Fulgent Genetics   |
| <b>78CPG</b> | Early Onset Epileptic Encephalopathy NGS Panel (Sequencing & Deletion/Duplication)                              | Fulgent Genetics   |
| <b>77TTG</b> | Early Onset Epileptic Encephalopathy NGS Panel (Sequencing Only)  | Fulgent Genetics   |
| <b>77GTG</b> | Epilepsy Advanced Sequencing and CNV Evaluation   | Athena Diagnostics Inc                                     |
| <b>78FNG</b> | Epilepsy Advanced Sequencing and CNV Evaluation - Epileptic Encephalopathy                                      | Athena Diagnostics Inc                                     |
| <b>77J4G</b> | Epilepsy Advanced Sequencing and CNV Evaluation - Generalized, Absence, Focal, Febrile and Myoclonic Epilepsies | Athena Diagnostics Inc                                     |
| <b>77HNG</b> | Epilepsy Advanced Sequencing and CNV Evaluation - Infantile Spasms  | Athena Diagnostics Inc                                     |
| <b>78FPG</b> | Epilepsy Advanced Sequencing and CNV Evaluation - Intellectual Disability                                       | Athena Diagnostics Inc                                     |
| <b>77HQG</b> | Epilepsy Advanced Sequencing and CNV Evaluation - Syndromic Disorders   | Athena Diagnostics Inc                                     |
| <b>7DP9G</b> | Epilepsy and Seizure Panel  | PreventionGenetics, part of Exact Sciences                 |
| <b>5DSFG</b> | Epilepsy Comprehensive NGS Panel (Deletion/Duplication Only)  | Fulgent Genetics   |
| <b>7724G</b> | Epilepsy Comprehensive NGS Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics   |
| <b>77D4G</b> | Epilepsy Comprehensive NGS Panel (Sequencing Only)  | Fulgent Genetics   |
| <b>6DDAG</b> | Epilepsy Deletion/Duplication Panel   | GeneDx   |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>  |
|--------------|--|---|
| <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>78FKG</b> | EPILEPSY GENE PANEL, COMPREHENSIVE WHEN THERAPEUTIC PANEL NEGATIVE | Ann and Robert Lurie Children's Hospital of Chicago                             |
| <b>77HJG</b> | EPILEPSY GENE PANEL, INFANTILE                                     | Ann and Robert Lurie Children's Hospital of Chicago                             |
| <b>34NLG</b> | EPILEPSY GENE PANEL, NOCTURNAL FRONTAL LOBE                        | Ann and Robert Lurie Children's Hospital of Chicago                             |
| <b>77YAG</b> | EPILEPSY GENE PANEL, THERAPEUTIC                                   | Ann and Robert Lurie Children's Hospital of Chicago                             |
| <b>7V2RG</b> | Epilepsy Panel   | Baylor Genetics, LLC  |
| <b>78FEG</b> | Epilepsy Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics           |
| <b>7XHKG</b> | Epilepsy Panel   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| <b>79Q4G</b> | Epilepsy panel - NGS Panel (CNV included)                          | Centogene   |
| <b>77Y5G</b> | Epilepsy/Seizure NGS Panel   | Greenwood Genetic Center - Molecular Diagnostic Laboratory                      |
| <b>77LQG</b> | Epilepsy/Seizures panel  | Knight Molecular Diagnostic   |
| <b>78G2G</b> | EpilepsyNext   | Ambry Genetics  |
| <b>78GSG</b> | EpilepsyNext-Expanded  | Ambry Genetics  |
| <b>776GG</b> | Epileptic Encephalopathy (NGS Panel and Copy Number Analysis)      | MNG Laboratories  |
| <b>86PAG</b> | EpiPanelDxPLUS   | Bionano Laboratories  |
| <b>772UG</b> | EpiRapid   | Ambry Genetics  |
| <b>775EG</b> | EpiXpanded Panel   | GeneDx  |
| <b>6YMKG</b> | Genomic Unity Epilepsy Analysis                                    | Varietyx, Inc.  |
| <b>77P9G</b> | Infantile Epilepsy Panel   | Knight Molecular Diagnostic   |
| <b>787MG</b> | Infantile Epilepsy Panel   | PerkinElmer Genomics  |
| <b>3Q7DG</b> | Infantile Spasms NGS Panel (Deletion/Duplication Only)             | Fulgent Genetics  |
| <b>76XZG</b> | Infantile Spasms NGS Panel (Sequencing & Deletion/Duplication)     | Fulgent Genetics  |

| GTU   | Test Name   | Laboratory Name                            |
|-------|---|--|
| 76WJG | Infantile Spasms NGS Panel (Sequencing Only)                                      | Fulgent Genetics                           |
| 76TEG | Invitae Epilepsy Panel  | Invitae Corporation                        |
| 7WZ6G | Invitae Epilepsy Panel - UNLOCKBTS  | Invitae Corporation                        |
| 7834G | Invitae Epilepsy Panel-Add-on Preliminary-evidence Genes for Epilepsy             | Invitae Corporation                        |
| 7WZ7G | Invitae Epilepsy Panel-Add-on Preliminary-evidence Genes for Epilepsy - UNLOCKBTS | Invitae Corporation                        |
| 5DQXG | Neonatal Epilepsy NGS Panel (Deletion/Duplication Only)                           | Fulgent Genetics                           |
| 77VEG | Neonatal Epilepsy NGS Panel (Sequencing & Deletion/Duplication)                   | Fulgent Genetics                           |
| 77VNG | Neonatal Epilepsy NGS Panel (Sequencing Only)                                     | Fulgent Genetics                           |
| 78HSG | PGmax™ - Comprehensive Epilepsy and Seizure Panel                                 | PreventionGenetics, part of Exact Sciences |
| 77XKG | PGXome Custom - Early Epileptic Encephalopathy, Dominant and X-linked             | PreventionGenetics, part of Exact Sciences |
| 7764G | PGXome Custom - Early Infantile Epileptic Encephalopathy, Dominant and X-linked   | PreventionGenetics, part of Exact Sciences |
| 78EUG | PGXome Custom - Early Infantile Epileptic Encephalopathy, Recessive               | PreventionGenetics, part of Exact Sciences |
| 7B4AG | STAT Comprehensive Epilepsy Panel   | PerkinElmer Genomics                       |
| 7V2QG | STAT Epilepsy Panel   | Baylor Genetics, LLC                       |
| 7AY4G | STAT Infantile Epilepsy Panel   | PerkinElmer Genomics                       |

### Evidence-Based Lung Cancer Diagnostic Algorithmic Tests

| GTU   | Test Name                                  | Laboratory Name |
|-------|--|-----------------|
| 7RVVG | Nodify Lung Nodule Risk Assessment Testing | Biodesix        |
| 6USVG | Nodify XL2                                 | Biodesix        |

## Exome Sequencing Tests

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 7AY2G | Add Familial Report to Previous Whole Exome Sequencing TRIO Test                   | PerkinElmer Genomics  |
| 79AQG | CentoDx - NGS Panel  | Centogene   |
| 79ARG | CentoLCV - NGS Panel (CNV included)  | Centogene   |
| 46YFG | CHOP Medical Exome   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 46WXG | CHOP Medical Exome + MitoGenome Combined Test                                      | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 6RG2G | Chromosomal Sequencing Analysis (Sequencing & Deletion/Duplication)                | Fulgent Genetics  |
| 77JAG | Clinical Exome (Sequencing & Del/Dup) Trio   | Fulgent Genetics  |
| 77YSG | Clinical Exome (Sequencing & Deletion/Duplication)                                 | Fulgent Genetics  |
| 78GBG | Clinical Exome (Sequencing Only)   | Fulgent Genetics  |
| 78GCG | Clinical Exome (Sequencing) Trio   | Fulgent Genetics  |
| 7BMXG | Clinical Exome (Sequencing) Trio   | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 24C9G | Clinical Exome Sequencing - Duo (Proband and 1 family member)                      | UCLA Diagnostic Molecular Pathology Laboratory                        |
| 2WCZG | Clinical Exome Sequencing - Exome re-analysis                                      | UCLA Diagnostic Molecular Pathology Laboratory                        |
| 33B2G | Clinical Exome Sequencing - Family Trio (Proband and two family members preferred) | UCLA Diagnostic Molecular Pathology Laboratory                        |
| 2XH2G | Clinical Exome Sequencing - Individual/Proband only                                | UCLA Diagnostic Molecular Pathology Laboratory                        |
| 2ZN9G | Clinical Exome Sequencing - Quad (Proband and 3 family members)                    | UCLA Diagnostic Molecular Pathology Laboratory                        |
| 7BMKG | Clinical Exome Sequencing (Duo)  | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 7BMVG | Clinical Exome Sequencing (Proband Only)   | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 7BMWG | Clinical Exome Sequencing (Quad)   | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 7TYRG | Critical Trio Whole Exome Sequencing   | University of Michigan - Michigan Medical Genetics Laboratories       |
| 7VBYG | Duo Exome  | AiLife Diagnostics  |
| 86SUG | Duo Whole Exome Sequencing   | Baylor Genetics, LLC  |
| 3V27G | Exome Select   | The University of Chicago Genetic Services                            |
| 7UNTG | Exome Sequencing   | ARUP Laboratories   |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>   |
|--------------|--|--|
| <b>7VSBG</b> | Exome Sequencing   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory                |
| <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>7VSAG</b> | Exome Sequencing Result  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory                |
| <b>53RNG</b> | Exome Sequencing Symptom-Guided Analysis                       | UCSD - Molecular Diagnostics & Cytogenetics Laboratory   |
| <b>4N3SG</b> | Exome Sequencing Trio  | Northwest Clinical Genomics Lab  |
| <b>7UNSG</b> | Exome Sequencing, Familial Control                             | ARUP Laboratories  |
| <b>46SRG</b> | Exome, Family Member   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics                            |
| <b>2YZMG</b> | ExomeNext-Duo  | Ambry Genetics   |
| <b>28FHG</b> | ExomeNext-Duo plus mtDNA                                       | Ambry Genetics   |
| <b>2YZRG</b> | ExomeNext-Proband  | Ambry Genetics   |
| <b>2YZQG</b> | ExomeNext-Proband plus mtDNA                                   | Ambry Genetics   |
| <b>2YZVG</b> | ExomeNext-Trio   | Ambry Genetics   |
| <b>5U8XG</b> | ExomeNext-Trio plus mtDNA                                      | Ambry Genetics   |
| <b>7XW6G</b> | ExomeReveal™   | Ambry Genetics   |
| <b>33Z3G</b> | ExomeSeq - Whole Exome Sequencing                              | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| <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>7A7AG</b> | Family Member Comparator Specimen for Exome Sequencing, Varies | Mayo Clinic Laboratories   |

| <b>GTU</b>   | <b>Test Name</b>  | <b>Laboratory Name</b>   |
|--------------|---|--|
| <b>7WZQG</b> | Family Targeted Testing   | AiLife Diagnostics   |
| <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>6ZSHG</b> | Genomic Unity Exome Analysis  | Varietyx, Inc.   |
| <b>7ABTG</b> | Genomic Unity Exome Plus Analysis                                     | Varietyx, Inc.   |
| <b>2BFBG</b> | GML Exome Family Member Peripheral blood draw                         | UCSF Molecular Diagnostics Laboratory  |
| <b>7WADG</b> | Hereditary Exome Family Member  | University of Minnesota Physicians Outreach Laboratory   |
| <b>7W4ZG</b> | Hereditary Whole Exome Sequencing                                     | University of Minnesota Physicians Outreach 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>6RLPG</b> | MNG Exome Proband Only Sequencing                                     | MNG Laboratories   |
| <b>6RLJG</b> | MNG Exome Proband Only Sequencing + mtDNA                             | MNG Laboratories   |
| <b>6RGKG</b> | MNG Exome TRIO Sequencing   | MNG Laboratories   |
| <b>77FXG</b> | MNG Exome TRIO Sequencing + mtDNA                                     | MNG Laboratories   |
| <b>4AVGG</b> | NextStepDx PLUS   | Bionano Laboratories   |
| <b>5E4FG</b> | PGxome Diagnostic Exome Test - Duo                                    | PreventionGenetics, part of Exact Sciences   |
| <b>5E5HG</b> | PGxome Diagnostic Exome Test - Trio                                   | PreventionGenetics, part of Exact Sciences   |
| <b>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>5E3RG</b> | PGxome Diagnostic   | PreventionGenetics, part of Exact Sciences   |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>   |
|--------------|--|--|
| <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>7VBUG</b> | Proband Exome  | AiLife Diagnostics   |
| <b>2ZBFG</b> | Proband Whole Exome Sequencing   | Baylor Genetics, LLC   |
| <b>27YXG</b> | Proband Whole Exome Sequencing + Chromosomal Microarray Analysis (CMA) (Comprehensive) | Baylor Genetics, LLC   |
| <b>76VCG</b> | Proband Whole Exome Sequencing + Comprehensive mtDNA Analysis                          | Baylor Genetics, LLC   |
| <b>7XAXG</b> | Rapid Whole Exome Sequencing Patient Only  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                  |
| <b>2BJCG</b> | Reanalysis and Interpretation for WES (Internal)                                       | PerkinElmer Genomics   |
| <b>2LFVG</b> | REFLEX to Exome Sequencing   | Northwest Clinical Genomics Lab  |
| <b>6DM5G</b> | Reflex to Whole Exome after Slice (Proband Only)                                       | GeneDx   |
| <b>365BG</b> | Reflex to Whole Exome after Slice (Trio)   | GeneDx   |
| <b>7B6CG</b> | Reflex to Whole Exome Sequencing (from panel)  | PerkinElmer Genomics   |
| <b>2G7GG</b> | Sequential Trio Whole Exome Sequencing   | Baylor Genetics, LLC   |
| <b>7X4AG</b> | Somatic Disease/Germline Comparator Exome  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                  |
| <b>348XG</b> | Total BluePrint Panel  | Baylor Genetics, LLC   |
| <b>7VBWG</b> | Trio Exome   | AiLife Diagnostics   |
| <b>6QQMG</b> | Trio Whole Exome Sequencing  | Baylor Genetics, LLC   |
| <b>776NG</b> | Trio Whole Exome Sequencing + Comprehensive mtDNA Analysis                             | Baylor Genetics, LLC   |
| <b>2BFEG</b> | UCSF Genomics Blood Draw   | UCSF Molecular Diagnostics Laboratory  |
| <b>2LNHG</b> | WES - Additional Affected Sibling  | Baylor Genetics, LLC   |
| <b>22DJG</b> | WES Single or Multi-Sample (Duo, Trio, Quad, etc.)                                     | Medical College of Wisconsin - Human and Molecular Genetics Center                               |
| <b>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   |



| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 3AALG | Whole Exome (Sequencing) Trio   | Fulgent Genetics  |
| 7DQDG | Whole Exome and Mitochondrial Genome Sequencing, Varies               | Mayo Clinic Laboratories  |
| 2Z7LG | Whole Exome Sequencing  | Gene by Gene  |
| 6QZFG | Whole Exome Sequencing  | Greenwood Genetic Center - Molecular Diagnostic Laboratory                      |
| 7WZRG | Whole Exome Sequencing  | Otogenetics   |
| 7RSXG | Whole Exome Sequencing  | Rady Children's Institute for Genomic Medicine                                  |
| 7R8CG | Whole Exome Sequencing  | Washington University in St. Louis Genomics and Pathology Services              |
| 5JQPG | Whole Exome Sequencing - DUO (Proband)                                | LabCorp   |
| 7PQEG | Whole Exome Sequencing – DUO (Proband), Products of Conception (POC)  | LabCorp   |
| 7X3VG | Whole Exome Sequencing - Parental Sample                              | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 5JR8G | Whole Exome Sequencing - Proband Only                                 | LabCorp   |
| 7PQFG | Whole Exome Sequencing – Proband Only, Products of Conception (POC)   | LabCorp   |
| 7X4BG | Whole Exome Sequencing - Reanalysis                                   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 5XKLG | Whole Exome Sequencing - TRIO (Proband)                               | LabCorp   |
| 7PQDG | Whole Exome Sequencing – TRIO (Proband), Products of Conception (POC) | LabCorp   |
| 6LSZG | Whole Exome Sequencing Comparator - Additional FM                     | LabCorp   |
| 6LT2G | Whole Exome Sequencing Comparator - Father                            | LabCorp   |
| 6LSYG | Whole Exome Sequencing Comparator - Mother                            | LabCorp   |
| 7A7EG | Whole Exome Sequencing for Hereditary Disorders , Varies              | Mayo Clinic Laboratories  |
| 3SMCG | Whole Exome Sequencing Proband Only                                   | PerkinElmer Genomics  |
| 7AYYG | Whole Exome Sequencing QUAD   | PerkinElmer Genomics  |
| 7VXCG | Whole Exome Sequencing Reflex   | Baylor Genetics, LLC  |
| 2DR8G | Whole Exome Sequencing TRIO   | PerkinElmer Genomics  |
| 7AYLG | Whole Exome Sequencing, DUO   | PerkinElmer Genomics  |
| 2D8LG | Whole Exome Sequencing: Duo Analysis                                  | University of Wisconsin - Madison WSLH - UW Cytogenetic Services                |

| <b>GTU</b>    | <b>Test Name</b>   | <b>Laboratory Name</b>  |
|---------------|--|---|
| <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>5FMPPG</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>7WZHG</b>  | Whole Exome with RISE Analysis (RNA-Seq) (Sequencing & Deletion/Duplication) | Fulgent Genetics  |
| <b>7TLFG</b>  | Whole Exome, 4 Sample  | Praxis Genomics, LLC  |
| <b>7TLDG</b>  | Whole Exome, Duo   | Praxis Genomics, LLC  |
| <b>7TLCG</b>  | Whole Exome, Proband   | Praxis Genomics, LLC  |
| <b>7TLEG</b>  | Whole Exome, Trio  | Praxis Genomics, LLC  |
| <b>6L9SG</b>  | XomeDx - Proband   | GeneDx  |
| <b>6RVYG</b>  | XomeDx - Trio  | GeneDx  |
| <b>6L9QG</b>  | XomeDx - Duo   | GeneDx  |
| <b>7TVKG</b>  | XomeDx Express (GeneDx), Blood   | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>7TVLG</b>  | XomeDx Express, Buccal Swab  | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>4JJWG</b>  | XomeDx Fetal - Duo   | GeneDx  |
| <b>7DQWG</b>  | XomeDx Plus - Duo  | GeneDx  |
| <b>77MHG</b>  | XomeDx Plus - Proband  | GeneDx  |
| <b>77M2G</b>  | XomeDx Plus - Trio   | GeneDx  |
| <b>7TV8G</b>  | XomeDx Plus- Duo, blood  | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>7TV7G</b>  | XomeDx Plus- Duo, buccal kit   | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>7TVBG</b>  | XomeDx Plus- Proband, blood  | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>7TV9G</b>  | XomeDx Plus- Trio, blood   | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>7TVCG</b>  | XomeDX Plus-Trio, buccal kit   | University of Michigan - Michigan Medical Genetics Laboratories       |
| <b>2W48G</b>  | XomeDx Prenatal - Comprehensive  | GeneDx  |
| <b>46N6G</b>  | XomeDx Trio  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |

| GTU   | Test Name            | Laboratory Name   |
|-------|----------------------|---|
| 4HM8G | XomeDxFetal- Proband | GeneDx  |
| 4GZRG | XomeDxFetal- Trio    | GeneDx  |
| 46J4G | XomeDxPlus Proband   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 46V3G | XomeDxPlus Trio      | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |

### Expanded Carrier Panel Tests

| GTU   | Test Name  | Laboratory Name  |
|-------|--|------------------|
| 7SSBG | Beacon 787-Expanded Carrier Screening Panel (With X-linked Disorders) (Sequencing & Deletion/Duplication)                          | Fulgent Genetics |
| 7SSCG | Beacon 787-Expanded Carrier Screening Panel (Without X-linked Disorders) (Sequencing & Deletion/Duplication)                       | Fulgent Genetics |
| 86GSG | Beacon ACMG Tier 3 Female Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 86GTG | Beacon ACMG Tier 3 Male Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 6LMNG | Beacon ACOG/ACMG Female Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 5DRBG | Beacon ACOG/ACMG Male Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 78HAG | Beacon Expanded Female Carrier Screening Panel (Sequencing & Deletion/Duplication)   | Fulgent Genetics |
| 77ZHG | Beacon Expanded Female Carrier Screening Plus Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 77ZJG | Beacon Expanded Male Carrier Screening Panel (Sequencing & Deletion/Duplication)   | Fulgent Genetics |
| 7743G | Beacon Expanded Male Carrier Screening Plus Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 77K3G | Beacon Focus Female Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 78HBG | Beacon Focus Male Carrier Screening Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics |
| 7WZBG | Beacon Preconception 787-Expanded Carrier Screening Panel (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication) | Fulgent Genetics |

| GTU   | Test Name   | Laboratory Name                       |
|-------|---|---------------------------------------|
| 7WZCG | Beacon Preconception 787-Expanded Carrier Screening Panel (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication) | Fulgent Genetics                      |
| 7WZDG | Beacon Preconception ACMG Tier 3 Carrier Screening (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication)             | Fulgent Genetics                      |
| 7WZEG | Beacon Preconception ACMG Tier 3 Carrier Screening (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication)        | Fulgent Genetics                      |
| 7WZ9G | Beacon Preconception Expanded Carrier Screening (With X-linked Disorders and Opt-In) (Sequencing & Deletion/Duplication)                | Fulgent Genetics                      |
| 7WZAG | Beacon Preconception Expanded Carrier Screening (Without X-linked Disorders; With Opt-In) (Sequencing & Deletion/Duplication)           | Fulgent Genetics                      |
| 7QH4G | CARRIER SCREEN - CLASSIC, FEMALE (24 GENE PANEL)  | Clinical Pathology Laboratories       |
| 7QH5G | CARRIER SCREEN - CLASSIC, MALE (22 GENE PANEL)  | Clinical Pathology Laboratories       |
| 7QGZG | CARRIER SCREEN - COMPLETE, MALE (147 GENE PANEL)  | Clinical Pathology Laboratories       |
| 79SAG | Carrier Screening (Horizon)   | ProPath                               |
| 6U8PG | Common Carrier Screening Panel  | Connective Tissue Gene Tests          |
| 7V3RG | Comprehensive Carrier Screening Panel for Genetic Conditions  | Genesys Diagnostics Inc               |
| 78GTG | Expanded Carrier Screen by Next Generation Sequencing with Fragile X  | ARUP Laboratories                     |
| 7832G | Expanded Carrier Screening  | UCSF Molecular Diagnostics Laboratory |
| 86D7G | Extended Carrier Screening Panel  | Connective Tissue Gene Tests          |
| 6RN4G | Foresight Fundamental Plus panel  | Myriad Genetics                       |
| 78HYG | Foresight Universal Panel Carrier Screen  | Myriad Genetics                       |
| 34P9G | FUNDAMENTAL PLUS 2528 (COUNSYL)   | Enzo Clinical Labs                    |
| 7V27G | GeneAware - Complete Panel - Female   | Baylor Genetics, LLC                  |
| 7V22G | GeneAware - Complete Panel - Male   | Baylor Genetics, LLC                  |
| 7V26G | GeneAware™ ACMG & ACOG Panel (Female)   | Baylor Genetics, LLC                  |
| 7UZZG | GeneAware™ ACMG & ACOG Panel (Male)   | Baylor Genetics, LLC                  |
| 7V24G | GeneAware™ Expanded Panel (Female)  | Baylor Genetics, LLC                  |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>  |
|--------------|--|---|
| <b>7UZYG</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>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>7X2CG</b> | GxVISION Carrier Screening Test ACOG/ACMG with CF            | Otogenetics   |
| <b>7X29G</b> | GxVISION Carrier Screening Test Pan-Ethnic Carrier Screening | Otogenetics   |
| <b>7759G</b> | Horizon 14 (14 disease panel)                                | Natera  |
| <b>7758G</b> | Horizon 27 (27 disease Pan-ethnic Standard panel)            | Natera  |
| <b>7757G</b> | Horizon 274 (274 disease Pan-ethnic Extended panel)          | Natera  |
| <b>72ESG</b> | Horizon 421  | Natera  |
| <b>7A7WG</b> | Horizon ACMG Panel   | Natera  |
| <b>6V4QG</b> | INHERIGEN  | GenPath Diagnostics   |
| <b>6UWPG</b> | INHERIGEN PLUS   | GenPath Diagnostics   |
| <b>6Y5WG</b> | InheriGenTx  | BioReference Laboratories   |
| <b>7SUSG</b> | Inheritest 100 PLUS Panel                                    | LabCorp   |
| <b>7SUTG</b> | Inheritest 300 PLUS Panel                                    | LabCorp   |
| <b>77LMG</b> | Inheritest 500 PLUS Panel                                    | Integrated Genetics   |
| <b>77BTG</b> | Inheritest 500 PLUS Panel                                    | LabCorp   |
| <b>7SURG</b> | Inheritest High Frequency Panel                              | LabCorp   |
| <b>7838G</b> | Inheritest 500 PLUS with Repro Partners Report               | Integrated Genetics   |
| <b>77ENG</b> | Inheritest Carrier Screen - Society Guided Panel (14 Genes)  | Integrated Genetics   |
| <b>7SRSG</b> | M Beacon Focus B, 14 Gene Panel                              | LabCorp   |
| <b>7ZPEG</b> | MYRIAD FORESIGHT CARRIER SCREENING                           | University of Illinois at Chicago - Biochemical Genetics Laboratory |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 784RG | MYRIAD WOMEN\\S HEALTH FAMILY PREP SCREEN 2            | University of California Davis Health System - UCDMC - Molecular and Cytogenetic Laboratory |
| 78NPG | Natera One Panel                                       | ProPath   |
| 3EHWG | NewbornGenelD  | Advanced Molecular Diagnostics, LLC   |
| 2D58G | NewbornGenelD include Fragile X (Female Patients Only) | Advanced Molecular Diagnostics, LLC   |
| 7SUAG | QHerit; 381 Diseases, Male                             | Quest Diagnostics   |
| 7VXWG | QHerit; 400 Diseases, Male                             | Quest Diagnostics   |
| 7SU9G | QHerit; 421 Diseases, Female                           | Quest Diagnostics   |
| 7VXXG | QHerit; 445 Diseases, Female                           | Quest Diagnostics   |
| 6UVXG | QHerit; Expanded Carrier Screen                        | Quest Diagnostics   |
| 7DHGG | QHerit; Extended, Female                               | Quest Diagnostics   |
| 7DHHG | QHerit; Extended, Male                                 | Quest Diagnostics   |
| 7DHJG | QHerit; Plus, Female                                   | Quest Diagnostics   |
| 7DHKG | QHerit; Plus, Male                                     | Quest Diagnostics   |
| 4TGTG | UNIVERSAL GENETIC TEST 2521 (COUNSYL)                  | Enzo Clinical Labs  |
| 4B77G | UNIVERSAL PANEL PLUS THROMBOPHILIAS 2524 (COUNSYL)     | Enzo Clinical Labs  |
| 4CK8G | UNIVERSAL PLUS GENETICS TEST 2522 (COUNSYL)            | Enzo Clinical Labs  |

### Genome Reanalysis Tests (Interpretation Only)

| GTU   | Test Name  | Laboratory Name      |
|-------|--|----------------------|
| 7B8KG | Analysis and Interpretation of Whole Genome Sequencing External Data | PerkinElmer Genomics |
| 7TL3G | External Data Analysis, 4 Sample                                     | Praxis Genomics, LLC |
| 7TKZG | External Data Analysis, Duo  | Praxis Genomics, LLC |
| 7TKYG | External Data Analysis, Proband                                      | Praxis Genomics, LLC |
| 7TL2G | External Data Analysis, Trio   | Praxis Genomics, LLC |
| 2LUBG | GenomeSeqDx 2nd Reanalysis   | GeneDx               |
| 2WWMG | GenomeSeqDx Reanalysis   | GeneDx               |

| GTU   | Test Name   | Laboratory Name          |
|-------|---|--------------------------|
| 7VP7G | Reanalysis and Interpretation of Previous RVTY Whole Genome DUO Sequencing Test     | PerkinElmer Genomics     |
| 7AXXG | Reanalysis and Interpretation of Previous RVTY Whole Genome Proband Sequencing Test | PerkinElmer Genomics     |
| 7VP6G | Reanalysis and Interpretation of Previous RVTY Whole Genome QUAD Sequencing Test    | PerkinElmer Genomics     |
| 7C4HG | Reanalysis and Interpretation of Previous RVTY Whole Genome TRIO Sequencing Test    | PerkinElmer Genomics     |
| 7VP8G | Reanalysis of Secondary Findings of Previous RVTY Test                              | PerkinElmer Genomics     |
| 7V2YG | WGS Reanalysis  | Baylor Genetics, LLC     |
| 7T4HG | Whole Genome Reanalysis   | ARUP Laboratories        |
| 7SAUG | Whole Genome Sequencing Reanalysis, Varies  | Mayo Clinic Laboratories |

## Genome Sequencing Tests

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 7AY3G | Add Familial Report to Previous Whole Genome Sequencing TRIO Test | PerkinElmer Genomics  |
| 7V9YG | Duo WGS   | AiLife Diagnostics  |
| 86SWG | Duo Whole Genome Sequencing                                       | Baylor Genetics, LLC  |
| 7SC9G | Family Member Comparator Specimen for Genome Sequencing, Varies   | Mayo Clinic Laboratories  |
| 7TVEG | Genome Express  | University of Michigan - Michigan Medical Genetics Laboratories                 |
| 7XC6G | Genome Sequencing-Parent Sample                                   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 7VRMG | GenomeSeqDx - Duo   | GeneDx  |
| 7VRNG | GenomeSeqDx - Proband   | GeneDx  |
| 6DX7G | GenomeSeqDx - Trio  | GeneDx  |
| 749YG | Genomic Unity® Whole Genome Analysis                              | Varietyx, Inc.  |
| 2NVZG | GenomX Sequencing Test  | Gene by Gene  |
| 86PCG | IriSight™? Comprehensive Analysis – Prenatal                      | Varietyx, Inc.  |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>  |
|--------------|--|---|
| <b>6R9PG</b> | MNGenome Additional Comparator (only available when trio is ordered) | MNG Laboratories  |
| <b>6R9SG</b> | MNGenome DUO Sequencing  | MNG Laboratories  |
| <b>6R9QG</b> | MNGenome Proband Only Sequencing                                     | MNG Laboratories  |
| <b>6RLTG</b> | MNGenome TRIO Sequencing   | MNG Laboratories  |
| <b>4E8TG</b> | PGnome Health Screen   | PreventionGenetics, part of Exact Sciences                                      |
| <b>6S6NG</b> | PGnome® Diagnostic - Duo   | PreventionGenetics, part of Exact Sciences                                      |
| <b>6S6QG</b> | PGnome® Diagnostic - Trio  | PreventionGenetics, part of Exact Sciences                                      |
| <b>6LTUG</b> | PGnome™ - 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>7XCAG</b> | Reanalysis Genome Sequencing-Parent Sample                           | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| <b>7RVSG</b> | Reflex to Whole Genome Sequencing (from genome panels)               | PerkinElmer Genomics  |
| <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>7WZGG</b> | Whole Exome with RISE Analysis (RNA-Seq) (Sequencing Only)           | Fulgent Genetics  |
| <b>7U3WG</b> | Whole Genome Sequencing  | ARUP Laboratories   |
| <b>6RPTG</b> | Whole Genome Sequencing  | Children's Mercy Hospital and Clinics - Molecular Genetics Laboratory           |
| <b>52SUG</b> | Whole Genome Sequencing  | Gene by Gene  |
| <b>27KEG</b> | Whole Genome Sequencing  | Johns Hopkins Medical Institutions - Pathology Laboratory                       |



| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 7TYTG | Whole Genome Sequencing (Baylor)  | University of Michigan - Michigan Medical Genetics Laboratories    |
| 7SAVG | Whole Genome Sequencing for Hereditary Disorders, Varies                | Mayo Clinic Laboratories   |
| 3VU6G | Whole Genome Sequencing Proband Only                                    | PerkinElmer Genomics   |
| 22DHG | Whole Genome Sequencing Single and Multi-Family (Duo, Trio, Quad, etc.) | Medical College of Wisconsin - Human and Molecular Genetics Center |
| 7TKXG | Whole Genome Sequencing, 4 Sample                                       | Praxis Genomics, LLC   |
| 7B2AG | Whole Genome Sequencing, DUO  | PerkinElmer Genomics   |
| 7TKVG | Whole Genome Sequencing, DUO  | Praxis Genomics, LLC   |
| 7U3XG | Whole Genome Sequencing, Familial Control                               | ARUP Laboratories  |
| 7TKUG | Whole Genome Sequencing, Proband  | Praxis Genomics, LLC   |
| 7B2LG | Whole Genome Sequencing, QUAD   | PerkinElmer Genomics   |
| 2UAVG | Whole Genome Sequencing, TRIO   | PerkinElmer Genomics   |
| 7TKWG | Whole Genome Sequencing, TRIO   | Praxis Genomics, LLC   |
| 7TLHG | Whole Genome Upgrade per Sample   | Praxis Genomics, LLC   |
| 7TLNG | Whole Genome Upgrade per Sample   | Praxis Genomics, LLC   |

### Hematologic Malignancy Panel Tests

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 2FKWG | Comprehensive Hematologic Cancer Panel                 | Children's Hospital of Philadelphia - Division of Genomic Diagnostics                            |
| 7SFFG | Comprehensive HemeComplete Profile                     | PathGroup  |
| 2DVSG | Comprehensive Lymphoid Oncology (DNA and RNA analysis) | Children's Hospital Colorado Precision Diagnostics Laboratory                                    |
| 7SFEG | Comprehensive Myeloid Profile                          | PathGroup  |
| 5M87G | FISH Panel : Myeloid Disorders                         | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 7VMHG | FISH Panel: Myeloid Malignancy                         | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 6Z8DG | FoundationOne Heme                                     | Foundation Medicine  |
| 34YSG | GeneTrails Hematologic Malignancies 220 Gene Panel     | Knight Molecular Diagnostic  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 7XE7G | Hematologic Cancer Fusion Analysis  | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory   |
| 45EWG | Hematologic Cancer Panel  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| 73H3G | Hematologic Neoplasm Mutation Panel   | Ohio State University - Molecular Pathology Laboratory                            |
| 34PNG | HemaVision Leukemia Panel   | University of Oklahoma Health Sciences Center - Molecular Pathology Laboratory    |
| 7XRGG | Heme Amplicon Panel by NGS  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 7ATGG | HEME DNA MUTATION PANEL (152 gene DNA NGS panel for mutations)                            | Texas Children's Hospital   |
| 7ATHG | HEME DNA/RNA COMBINED PANEL (Combined heme mutation and fusion panels)                    | Texas Children's Hospital   |
| 6LTNG | Heme Gene Panel by NGS  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 7SFCG | HemeComplete NextGen Sequencing Assay   | PathGroup   |
| 72AHG | IntelliGEN Myeloid  | LabCorp   |
| 7RJGG | JMML Associated Exon Panel (JMML)   | Rady Children's Institute for Genomic Medicine                                    |
| 5PPVG | Juvenile Myelomonocytic Leukemia Associated Exon Panel                                    | UCSF Molecular Diagnostics Laboratory   |
| 28E8G | Leukemia Trial NGS Panel, Blood   | Johns Hopkins Medical Institutions - Pathology Laboratory                         |
| 28MEG | Leukemia Trial NGS Panel, Bone Marrow   | Johns Hopkins Medical Institutions - Pathology Laboratory                         |
| 6LTPG | Lymphoid Gene Panel by NGS  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 2DJPG | Lymphoid Oncology DNA Analysis  | Children's Hospital Colorado Precision Diagnostics Laboratory                     |
| 7XR9G | MayoComplete Chronic Lymphoid Neoplasms, Next-Generation Sequencing, Varies               | Mayo Clinic Laboratories  |
| 7XR8G | MayoComplete Histiocytic Neoplasms, Next-Generation Sequencing, Varies                    | Mayo Clinic Laboratories  |
| 6U9NG | MayoComplete Myeloid Neoplasms, Comprehensive OncoHeme Next-Generation Sequencing, Varies | Mayo Clinic Laboratories  |
| 7SBCG | MayoComplete Plasma Cell Myeloma, Next-Generation Sequencing, Varies                      | Mayo Clinic Laboratories  |
| 7XR7G | MayoComplete T-Cell Lymphoma, Next-Generation Sequencing, Varies                          | Mayo Clinic Laboratories  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 2XDBG | Myeloid Complete Molecular Profile  | Genetic Associates  |
| 6VB9G | Myeloid Extended Mutation Analysis Panel by Next Generation Sequencing                      | Molecular Pathology Laboratory Network  |
| 6LTQG | Myeloid Gene Panel by NGS   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory       |
| 7UXCG | Myeloid Malignancies Mutation and Copy Number Variation Panel by Next Generation Sequencing | ARUP Laboratories   |
| 7ALDG | MYELOID MALIGNANCIES MUTATION PANEL   | Clinical Pathology Laboratories   |
| 2XRDG | Myeloid Malignancies Mutation Panel by Next Generation Sequencing                           | ACL Laboratories  |
| 6VAZG | Myeloid Malignancies Mutation Panel by Next Generation Sequencing                           | ARUP Laboratories   |
| 78N9G | Myeloid Multigene Panel   | UCSF Molecular Diagnostics Laboratory   |
| 5CTBG | Myeloid Mutation Panel- AML   | University of North Carolina Hospitals - Molecular Genetics                             |
| 7Q2MG | Myeloid Mutation Panel, Blood   | Nebraska Medical Center - Molecular Diagnostic Laboratory                               |
| 7Q2HG | Myeloid Mutation Panel, Other   | Nebraska Medical Center - Molecular Diagnostic Laboratory                               |
| 8EFPG | Myeloid Neoplasm Mutation Analysis Only   | Medfusion   |
| 7SFBG | Myeloid NextGen Sequencing Assay  | PathGroup   |
| 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   |
| 6UY4G | Myeloid tumor panel - Somatic Mutation Analysis   | Centogene   |
| 7SPKG | Neo Comprehensive - Myeloid Disorders   | NeoGenomics Laboratories  |
| 7URQG | Neo Comprehensive™ - Heme Cancers   | NeoGenomics Laboratories  |
| 3ZSJG | NeoTYPE AITL/Peripheral T-Cell Lymphoma Profile   | NeoGenomics Laboratories  |
| 86UPG | NeoTYPE Lymphoid Disorders Profile  | NeoGenomics Laboratories  |
| 6VB7G | NGS Hematologic Malignancy Mutation Panel   | UCSD - Molecular Diagnostics & Cytogenetics Laboratory                                  |
| 2LJDG | NGS Hematolymphoid Panel (Lab Only)   | Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital |

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 5364G | NGS_Myeloid 37 Genes Panel                                       | CellNetix Pathology and Laboratories   |
| 86LPG | OnkoSight Advanced Chronic Lymphoid Neoplasm NGS Panel - PB/BM   | BioReference Laboratories  |
| 7C5AG | OnkoSight Advanced NGS Myeloid Panel                             | BioReference Laboratories  |
| 7T3WG | OnkoSight Advanced Pan Heme Fusion NGS Panel                     | BioReference Laboratories  |
| 7DN9G | Paired Tumor / Normal – Comprehensive Hematologic Cancer Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics          |
| 7DN8G | Paired Tumor / Normal - Hematologic Cancer Panel                 | Children's Hospital of Philadelphia - Division of Genomic Diagnostics          |
| 7UJHG | PennSeq Hematological Malignancies Panel                         | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory |
| 7W6YG | T Large Granular Lymphocyte NGS Panel                            | University of Minnesota Physicians Outreach Laboratory                         |
| 7ZQTG | Targeted Myeloid Panel (TMP)                                     | Columbia University - Personalized Genomic Medicine                            |
| 86SCG | Tempus  xT: Targeted panel of 648 genes (Hematologic Malignancy) | Tempus AI, Inc.  |

### Hereditary GI/Colon Cancer Panel Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 7TP3G | CancerNext Expanded Colon Lynch   | Ambry Genetics   |
| 7TNZG | CancerNext: Colon/Lynch   | Ambry Genetics   |
| 2NBHG | Centocolon - NGS Panel (CNV included)                                     | Centogene  |
| 76DFG | COLARIS AP PLUS with Myriad myRisk Hereditary Cancer Update Test          | Myriad Genetics  |
| 76DEG | COLARIS PLUS with Myriad myRisk Hereditary Cancer Update Test             | Myriad Genetics  |
| 76DUG | ColoNext  | Ambry Genetics   |
| 7WT3G | ColoNext  | University of Minnesota Physicians Outreach Laboratory |
| 6UQPG | ColoNext with RNAinsight  | Ambry Genetics   |
| 3598G | Colorectal Cancer Comprehensive Panel (Deletion/Duplication Only)         | Fulgent Genetics                                       |
| 76E3G | Colorectal Cancer Comprehensive Panel (Sequencing & Deletion/Duplication) | Fulgent Genetics                                       |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 3ATVG | Colorectal Cancer Comprehensive Panel (Sequencing Only)  | Fulgent Genetics  |
| 3D5AG | Colorectal Cancer Focus Panel (Deletion/Duplication Only)  | Fulgent Genetics  |
| 6U88G | Colorectal Cancer Focus Panel (Germline)   | NeoGenomics Laboratories  |
| 76E4G | Colorectal Cancer Focus Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics  |
| 34KAG | Colorectal Cancer Focus Panel (Sequencing Only)  | Fulgent Genetics  |
| 2L58G | Colorectal Cancer Germline NGS Panel   | University of Michigan - Michigan Medical Genetics Laboratories                   |
| 76DSG | Colorectal Cancer Panel  | ACL Laboratories  |
| 76DWG | Colorectal Cancer Panel  | GeneDx  |
| 76BGG | Colorectal Cancer Panel  | PerkinElmer Genomics  |
| 3DW5G | ColoSeq - Lynch and Polyposis Panel  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 6V2QG | Comprehensive Colon Cancer Panel (Seq & Del/Dup)   | Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories               |
| 7TNXG | Comprehensive Common Cancer Panel: Colon/Lynch   | GeneDx  |
| 25LZG | Hereditary Colorectal Cancer Panel   | Quest Diagnostics   |
| 76DZG | Hereditary Colorectal Cancer Panel   | The University of Chicago Genetic Services  |
| 7AAAG | Hereditary Colorectal Cancer Panel (20 Genes)  | Palo Verde Laboratory - division of Sonora Quest Lab                              |
| 7V2BG | Hereditary Colorectal/Gastrointestinal Cancer Panel  | Baylor Genetics, LLC  |
| 7SHSG | Hereditary Gastrointestinal Cancer High-Risk Panel, Sequencing and Deletion/Duplication                                    | ARUP Laboratories   |
| 76DNG | Hereditary Gastrointestinal Cancer Panel, Sequencing and Deletion/Duplication  | ARUP Laboratories   |
| 7DLGG | Hereditary Gastrointestinal Cancer Panel, Varies   | Mayo Clinic Laboratories  |
| 6V9EG | Hereditary High-Risk Colon Cancer Panel  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| 7V2GG | Hereditary High-Risk Colorectal Cancer Panel   | Baylor Genetics, LLC  |
| 45K7G | Inherited Colon Cancer Panel   | Knight Molecular Diagnostic   |
| 778DG | Invitae Hereditary Colorectal Cancer Guidelines-Based Panel  | Invitae Corporation   |
| 77QBG | Invitae Hereditary Colorectal Cancer Guidelines-Based Panel-Add-on Preliminary-evidence Colorectal Cancer Guidelines Genes | Invitae Corporation   |
| 76CYG | Invitae Hereditary Colorectal Cancer Panel   | Invitae Corporation   |

| GTU          | Test Name  | Laboratory Name      |
|--------------|--|----------------------|
| <b>76CZG</b> | Invitae Hereditary Colorectal Cancer Panel-Add-on Preliminary-evidence Genes for Colorectal Cancer | Invitae Corporation  |
| <b>6DXSG</b> | Lynch/Colorectal High Risk Panel   | GeneDx               |
| <b>7TNYG</b> | Rest of Comprehensive Common Cancer Panel: Colon/Lynch   | GeneDx               |
| <b>7B7KG</b> | STAT Colorectal Cancer Panel   | PerkinElmer Genomics |
| <b>7VPKG</b> | Tempus  xG Common Hereditary Cancers 36 genes (hereditary colon cancer indications)                | Tempus AI, Inc.      |
| <b>7VPGG</b> | Tempus  xG+ Extended Hereditary Cancers 88 genes (hereditary colon cancer indications)             | Tempus AI, Inc.      |
| <b>76BSG</b> | VistaSeq Colorectal Cancer Panel   | Integrated Genetics  |
| <b>76DAG</b> | VistaSeq Colorectal Cancer Panel   | LabCorp              |
| <b>76CBG</b> | VistaSeq Colorectal Cancer Panel   | Labcorp   Oncology   |
| <b>2F64G</b> | VistaSeq High Risk Colorectal Cancer Panel   | Integrated Genetics  |
| <b>58EYG</b> | VistaSeq High Risk Colorectal Cancer Panel   | LabCorp              |
| <b>4NLFG</b> | VistaSeq High Risk Colorectal Cancer Panel   | Labcorp   Oncology   |

### Hereditary Polyposis Panel Tests

| GTU          | Test Name  | Laboratory Name   |
|--------------|--|---|
| <b>7TQFG</b> | Adenomatous Polyposis Focus (Sequencing & Deletion/Duplication)                | Fulgent Genetics  |
| <b>5MJ8G</b> | APC & MUTYH seq and del/dup  | Ambry Genetics  |
| <b>6UQRG</b> | APC and MUTYH with RNAinsight  | Ambry Genetics  |
| <b>8734G</b> | APC- and MUTYH-Associated Polyposis Panel, Sequencing and Deletion/Duplication | ARUP Laboratories   |
| <b>2AUYG</b> | COLARIS AP   | Myriad Genetics   |
| <b>86HDG</b> | ColoSeq Genes Sequenced  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 86HCG | ColoSeq Polyposis   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 4QZBG | Comprehensive Polyposis Syndrome Panel (Seq & Del/Dup Analysis) | Genetics Center - Molecular and Cytogenetic Diagnostic Laboratories               |
| 4R4DG | FAP: APC Seq, APC Del/Dup, MUTYH 2 Muts                         | UCSD - Molecular Diagnostics & Cytogenetics Laboratory                            |
| 6QVMG | Hereditary Polyposis Panel                                      | PreventionGenetics, part of Exact Sciences  |
| 7PQHG | Invitae Adenomatous Polyposis Panel                             | Invitae Corporation   |
| 7TQGG | Polyposis Comprehensive (Sequencing & Deletion/Duplication)     | Fulgent Genetics  |

### Hypertrophic Cardiomyopathy (HCM) Panel Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 5LVCG | HCM Panel Deletion / Duplication  | DDC Clinic Laboratory  |
| 6PZ3G | HCM Panel Next Generation Sequencing                                      | DDC Clinic Laboratory  |
| 6PZ4G | HCM Panel NGS & Del Dup Comprehensive                                     | DDC Clinic Laboratory  |
| 2YZEG | HCMNext   | Ambry Genetics   |
| 2YZAG | HCMNext Reflex  | Ambry Genetics   |
| 5W3JG | Hypertrophic Cardiomyopathy – HCM (41 genes) Deletion/Duplication Panel   | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 2FPZG | Hypertrophic Cardiomyopathy – HCM (41 genes) NGS + Sanger fill-in Panel   | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 3XPUG | Hypertrophic Cardiomyopathy – HCM (41 genes) NGS Panel                    | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory                     |
| 3CEJG | Hypertrophic Cardiomyopathy (HCM) Panel                                   | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 7DN2G | Hypertrophic Cardiomyopathy (HCM) Panel                                   | Washington University in St. Louis Genomics and Pathology Services                               |
| 7SBZG | Hypertrophic Cardiomyopathy Gene Panel, Varies                            | Mayo Clinic Laboratories   |
| 5N9HG | Hypertrophic Cardiomyopathy NGS Panel                                     | Greenwood Genetic Center - Molecular Diagnostic Laboratory                                       |
| 3XG9G | Hypertrophic Cardiomyopathy NGS Panel (Deletion/Duplication Only)         | Fulgent Genetics   |
| 25E2G | Hypertrophic Cardiomyopathy NGS Panel (Sequencing & Deletion/Duplication) | Fulgent Genetics   |

| GTU    | Test Name   | Laboratory Name   |
|--------|---|---|
| 3XPSG  | Hypertrophic Cardiomyopathy NGS Panel (Sequencing Only)   | Fulgent Genetics  |
| 3XQ7G  | Hypertrophic Cardiomyopathy Panel   | Knight Molecular Diagnostic   |
| 7XH6G  | Hypertrophic Cardiomyopathy Panel   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory |
| 3XQBG  | Hypertrophic Cardiomyopathy Panel   | PerkinElmer Genomics  |
| 5CYCG  | Hypertrophic Cardiomyopathy Panel   | PreventionGenetics, part of Exact Sciences                                      |
| 3XNZG  | Hypertrophic Cardiomyopathy Panel (PCPGM)   | University of Michigan - Michigan Medical Genetics Laboratories                 |
| 3HV4G  | Hypertrophic Cardiomyopathy Panel (PCPGM), Known Mutation   | University of Michigan - Michigan Medical Genetics Laboratories                 |
| 2KTCCG | Hypertrophic Cardiomyopathy Panel, Sequencing   | ARUP Laboratories   |
| 44WDG  | Invitae Hypertrophic Cardiomyopathy Panel   | Invitae Corporation   |
| 42LUG  | Invitae Hypertrophic Cardiomyopathy Panel-Add-on Preliminary-evidence Genes for Hypertrophic Cardiomyopathy | Invitae Corporation   |
| 7B79G  | STAT Hypertrophic Cardiomyopathy Panel  | PerkinElmer Genomics  |

### Long QT Syndrome (LQTS) Panel Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 5FWSG | Invitae Long QT Syndrome Panel  | Invitae Corporation  |
| 786WG | Invitae Long QT Syndrome Panel-Add-on Preliminary-evidence Genes for Long QT Syndrome | Invitae Corporation  |
| 77MWG | Long and Short QT Syndrome (NGS Panel and Copy Number Analysis)                       | MNG Laboratories   |
| 6Q7RG | Long QT Panel Deletion / Duplication  | DDC Clinic Laboratory  |
| 782LG | Long QT Panel Next Generation Sequencing  | DDC Clinic Laboratory  |
| 782KG | Long QT Panel NGS & Del Dup Comprehensive   | DDC Clinic Laboratory  |
| 2KQNG | Long QT Panel, Sequencing and Deletion/Duplication                                    | ARUP Laboratories  |
| 39DQG | Long QT Syndrome – LQTS (15 genes) Deletion/Duplication Panel                         | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |
| 776UG | Long QT Syndrome – LQTS (15 genes) NGS + Sanger fill-in Panel                         | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |
| 78A2G | Long QT Syndrome – LQTS (15 genes) NGS Panel  | Baylor College of Medicine - John Welsh Cardiovascular Diagnostic Laboratory |



| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 77BKG | Long QT Syndrome (LQTS) Panel  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics   |
| 5XA8G | Long QT Syndrome (LQTS) Panel Sequencing   | Indiana University Molecular Genetics Diagnostic Laboratory - Indiana University Medical Center, School of Medicine |
| 27X3G | Long QT Syndrome and Short QT Syndrome   | Knight Molecular Diagnostic   |
| 77C6G | 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  |
| 7SBNG | Long QT Syndrome Gene Panel, Varies  | Mayo Clinic Laboratories  |
| 77L7G | Long QT syndrome NGS Panel   | Connective Tissue Gene Tests  |
| 5N7XG | Long QT syndrome NGS Panel   | Greenwood Genetic Center - Molecular Diagnostic Laboratory  |
| 6MADG | Long QT syndrome NGS Panel Deletion / Duplication  | Connective Tissue Gene Tests  |
| 774ZG | Long QT syndrome NGS Panel Next Generation Sequencing  | Connective Tissue Gene Tests  |
| 76V5G | Long QT Syndrome Panel   | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories                    |
| 7XHMG | Long QT Syndrome Panel   | Nationwide Children's Hospital - Cytogenetics and Molecular Genetics Laboratory                                     |
| 785ZG | Long QT Syndrome Panel   | PerkinElmer Genomics  |
| 78FTG | Long QT Syndrome Panel   | PreventionGenetics, part of Exact Sciences  |
| 7DN3G | Long QT Syndromes panel  | Washington University in St. Louis Genomics and Pathology Services  |
| 772GG | LongQTNext   | Ambry Genetics  |
| 783PG | LQTS Panel   | GeneDx  |
| 2YB5G | LQTS Seq/Del/Dup Panel   | Johns Hopkins Medical Institutions - Pathology Laboratory   |
| 7B7CG | STAT Long QT Syndrome Panel  | PerkinElmer Genomics  |

### Lung Cancer Panel Tests

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 8EEQG | 50SEQ with MSI Panel                                      | Medfusion  |
| 5X7JG | BRAF/EGFR NGS if EGFR neg rfx to ALK FISH if neg rfx ROS1 | BioReference Laboratories  |
| 2LVVG | Comprehensive Lung Panel                                  | University of Iowa Hospitals and Clinics - Department of Pathology |
| 7UQRG | Comprehensive LungSEQPlus Panel                           | Medfusion  |

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 7SNHG | Comprehensive OnkoSight Advanced Lung Panel                            | BioReference Laboratories  |
| 7TMTG | Comprehensive OnkoSight Lung Panel + GeneStrat Liquid Biopsy           | BioReference Laboratories  |
| 5XBBG | EGFR and KRAS, if both neg, reflex ALK, if neg reflex ROS1             | BioReference Laboratories  |
| 3XRVG | EGFR, KRAS, BRAF, PIK3CA mutation analysis, NSCLC panel                | University of Pittsburgh Medical Center - Division of Molecular Diagnostics                      |
| 4NUWG | FISH Panel : Non-Small Cell Lung Cancer                                | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 6LQSG | Lung adenocarcinoma targeted NGS panel                                 | Allina Health Laboratory   |
| 7AFUG | LUNG CANCER  | Clinical Pathology Laboratories  |
| 7Q8RG | Lung Cancer Mutation Analysis  | UCLA Diagnostic Molecular Pathology Laboratory   |
| 7VZQG | Lung Cancer Mutation Panel   | ARUP Laboratories  |
| 6WW7G | Lung Cancer Mutation Panel (EGFR, KRAS, ALK)                           | Quest Diagnostics  |
| 42L6G | Lung Cancer NGS Panel  | University of Michigan - Michigan Medical Genetics Laboratories                                  |
| 3ZETG | Lung Cancer Panel by Next Generation Sequencing                        | Molecular Pathology Laboratory Network   |
| 7R8HG | LungSEQ Concurrent Panel, without PD-L1                                | Quest Diagnostics  |
| 8EEXG | LUNGSEQ Panel  | Medfusion  |
| 7AUFG | LUNGSEQ Panel  | Quest Diagnostics  |
| 8EERG | LUNGSEQ Panel Concurrent   | Medfusion  |
| 8EFKG | LUNGSEQ Panel with FISH Concurrent - no PD-L1                          | Medfusion  |
| 7RWLG | MayoComplete Lung Cancer Mutations, Next-Generation Sequencing, Tumor  | Mayo Clinic Laboratories   |
| 7RWMG | MayoComplete Lung Cancer-Targeted Gene Panel with Rearrangement, Tumor | Mayo Clinic Laboratories   |
| 7RWKG | MayoComplete Lung Rearrangements, Rapid Test, Tumor                    | Mayo Clinic Laboratories   |
| 7BHXG | NeoTYPE® DNA & RNA - Lung  | NeoGenomics Laboratories   |
| 6YZRG | NeoTYPE® Lung Tumor Profile  | NeoGenomics Laboratories   |
| 262FG | Non-Small-cell Lung Cancer (NSCLC) Therapeutic Profile II              | LabCorp  |
| 6ST7G | Oncology FISH Analysis - Non-small Cell Lung Carcinoma Panel           | Baylor Genetics, LLC   |
| 86LVG | OnkoSight Advanced Comprehensive Lung                                  | BioReference Laboratories  |
| 86LLG | OnkoSight Advanced Lung Cancer NGS Panel                               | BioReference Laboratories  |

| GTU   | Test Name  | Laboratory Name                                      |
|-------|--|--|
| 6U25G | OnkoSight Advanced Lung Cancer Panel, ALK and ROS1 by FISH | BioReference Laboratories                            |
| 26QHG | Targeted Gene Panel with Fusions, Lung Cancer              | Palo Verde Laboratory - division of Sonora Quest Lab |

### Lung Cancer Treatment Algorithmic Tests

| GTU   | Test Name          | Laboratory Name |
|-------|--------------------|-----------------|
| 6XS9G | DetermaRX          | Oncocyte        |
| 7V98G | LungOI             | Imagene         |
| 7VN6G | PROphet NSCLC Test | OncoHost, Inc   |
| 6UT9G | VeriStrat          | Biodesix        |

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

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 6RPMG | COLARIS  | Myriad Genetics  |
| 7X23G | GxVISION Hereditary Cancer Risk Assessment Lynch Syndrome Genes  | Otogenetics  |
| 2YYWG | HNPCC concurrent   | Ambry Genetics   |
| 2FTJG | HNPCC PANEL  | Center for Human Genetics, Inc.                        |
| 7SNVG | HNPCC/Lynch Deletion/Duplication   | Ambry Genetics   |
| 2FS8G | HNPCC/Lynch Syndrome Del/Dup   | UCSD - Molecular Diagnostics & Cytogenetics Laboratory |
| 2AWXG | Invitae Lynch Syndrome Panel   | Invitae Corporation                                    |
| 2FSEG | Lynch Syndrome - Hereditary (Germline) Testing   | Labcorp   Oncology                                     |
| 7QFCG | Lynch Syndrome Focus (Sequencing & Deletion/Duplication)   | Fulgent Genetics                                       |
| 2FRUG | Lynch Syndrome Gene Panel: 5 Genes (EPCAM, MLH1, MSH2, MSH6, PMS2) by Gene Sequencing with Deletion/Duplication Analysis | Medical Diagnostic Laboratories, LLC                   |
| 7A2RG | Lynch syndrome Panel   | PerkinElmer Genomics                                   |
| 57DNG | Lynch syndrome Panel   | PreventionGenetics, part of Exact Sciences             |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 2FRZG | Lynch syndrome Panel   | Quest Diagnostics   |
| 5JURG | Lynch syndrome Panel   | The University of Chicago Genetic Services                      |
| 7AC3G | Lynch Syndrome Panel, Sequencing and Deletion/Duplication                  | ARUP Laboratories   |
| 7DL5G | Lynch Syndrome Panel, Varies   | Mayo Clinic Laboratories  |
| 4JKFG | Lynch Syndrome, MSH2 Sequencing and Deletion/Duplication (Including EPCAM) | Quest Diagnostics   |
| 72H6G | Lynch with RNAinsight  | Ambry Genetics  |
| 2FSKG | MLH1 / MSH2 / MSH6 / PMS2 Comprehensive Analysis                           | Labcorp   Oncology  |
| 2FSLG | MLH1 / MSH2 / MSH6 Comprehensive Analysis                                  | Labcorp   Oncology  |
| 2FSNG | MLH1 / MSH2 Comprehensive Analysis   | Labcorp   Oncology  |
| 7U2HG | MLH1, MSH2, MSH6, PMS2 Sequencing and Del/Dup (NGS)                        | University of Michigan - Michigan Medical Genetics Laboratories |
| 7DQLG | MLH1/MSH2 Del/Dup Testing by MLPA  | PerkinElmer Genomics  |
| 2YYNG | MSH2 seq & del/dup & EPCAM del/dup   | Ambry Genetics  |
| 7DQKG | MSH6/MUTYH/EPCAM Del/Dup Testing by MLPA                                   | PerkinElmer Genomics  |
| 87B9G | VistaSeq Lynch Syndrome Panel  | LabCorp   |

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

| GTU   | Test Name  | Laboratory Name                            |
|-------|--|--|
| 77W7G | Cellular Energetics Ophthalmoplegia Syndromes (NGS Panel and Copy Number Analysis + mtDNA) | MNG Laboratories                           |
| 5C94G | Chronic Progressive External Ophthalmoplegia (CPEO/PEO) Panel                              | PreventionGenetics, part of Exact Sciences |
| 6DD7G | Combined Mito Genome Plus Mito Focused Nuclear Gene Panel                                  | GeneDx                                     |
| 7R7VG | Combined Mitochondrial Full Genome and Nuclear Gene Panel, Varies                          | Mayo Clinic Laboratories                   |
| 6RKZG | Comprehensive Cellular Energetics Defects (NGS Panel and Copy Number Analysis + mtDNA)     | MNG Laboratories                           |
| 77WDG | Comprehensive Mitochondrial Nuclear Gene Panel   | PerkinElmer Genomics                       |
| 77SKG | Comprehensive mtDNA Depletion Syndromes (NGS Panel and Copy Number Analysis)               | MNG Laboratories                           |

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 76YBG | Comprehensive Ophthalmoplegia Syndromes (NGS Panel and Copy Number Analysis + mtDNA)         | MNG Laboratories   |
| 76YHG | Cytochrome C Oxidase Deficiency (NGS Panel and Copy Number Analysis + mtDNA)                 | MNG Laboratories   |
| 788KG | Dual Genome Leigh Disease Panel by Massively Parallel Sequencing                             | Baylor Genetics, LLC   |
| 6LC5G | Genomic Unity® Comprehensive Mitochondrial Disorders Analysis                                | Varietyx, Inc.   |
| 7V37G | Genomic Unity® Mitochondrial Genome Deletions Analysis                                       | Varietyx, Inc.   |
| 7V36G | Genomic Unity® Nuclear Encoded Mitochondrial Gene Analysis                                   | Varietyx, Inc.   |
| 3XEDG | Invitae Nuclear Mitochondrial Disorders Panel  | Invitae Corporation  |
| 77YFG | Leigh and Leigh-Like Syndrome Panel (Nuclear Genes Only)                                     | PreventionGenetics, part of Exact Sciences                                     |
| 2YAYG | Mito Disord Panel (mtDNA-108 Nuc Genes)  | UCSD - Molecular Diagnostics & Cytogenetics Laboratory                         |
| 77GLG | Mitochondrial Complex I Deficiency Panel (Nuclear Genes)                                     | PreventionGenetics, part of Exact Sciences                                     |
| 6QVJG | Mitochondrial Complex II Deficiency Panel  | PreventionGenetics, part of Exact Sciences                                     |
| 386QG | Mitochondrial Complex III Deficiency Panel (Nuclear Genes)                                   | PreventionGenetics, part of Exact Sciences                                     |
| 76ZNG | Mitochondrial Complex IV Deficiency Panel (Nuclear Genes)                                    | PreventionGenetics, part of Exact Sciences                                     |
| 5LQTG | Mitochondrial Complex V Deficiency Panel (Nuclear Genes)                                     | PreventionGenetics, part of Exact Sciences                                     |
| 7U5TG | Mitochondrial Disorders (mtDNA) Sequencing and Deletion Analysis by NGS                      | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory |
| 6LTZG | Mitochondrial Disorders Panel (Nuclear Genes Only)   | PreventionGenetics, part of Exact Sciences                                     |
| 6XJWG | Mitochondrial DNA Depletion Testing (Leukocyte)  | LabCorp  |
| 54EFG | Mitochondrial Encephalomyopathic Evaluation  | Athena Diagnostics Inc   |
| 77FTG | Mitochondrial Encephalopathy/Leigh Syndrome Nuclear Gene Panel                               | PerkinElmer Genomics   |
| 772TG | Mitochondrial Genome Maintenance/Integrity Nuclear Genes Panel                               | PreventionGenetics, part of Exact Sciences                                     |
| 3WQPG | Mitochondrial Hepatoencephalopathic Evaluation   | Athena Diagnostics Inc   |
| 4RDTG | Mitochondrial Respiratory Chain Complex II Deficiency Panel by Massively Parallel Sequencing | Baylor Genetics, LLC   |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>                     |
|--------------|--|--|
| <b>77T2G</b> | Mitochondrial Respiratory Chain Complex I-V Nuclear Gene Deficiency Panel by Massively Parallel Sequencing | Baylor Genetics, LLC                       |
| <b>62P9G</b> | mtDNA Depletion Syndrome NGS Panel (Deletion/Duplication Only)   | Fulgent Genetics                           |
| <b>77EVG</b> | mtDNA Depletion Syndrome NGS Panel (Sequencing & Deletion/Duplication)                                     | Fulgent Genetics                           |
| <b>77TJG</b> | mtDNA Depletion Syndrome NGS Panel (Sequencing Only)   | Fulgent Genetics                           |
| <b>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>776AG</b> | PGXome Custom - Combined Oxidative Phosphorylation Deficiency  | PreventionGenetics, part of Exact Sciences |
| <b>77KGG</b> | PGXome Custom - Comprehensive Cellular Energetics Defects  | PreventionGenetics, part of Exact Sciences |
| <b>77GJG</b> | PGXome Custom - Mitochondrial Complex I Deficiency   | PreventionGenetics, part of Exact Sciences |
| <b>4C5HG</b> | PGXome Custom - Mitochondrial Complex II Deficiency  | PreventionGenetics, part of Exact Sciences |
| <b>6P7LG</b> | PGXome Custom - Mitochondrial Complex III Deficiency   | PreventionGenetics, part of Exact Sciences |
| <b>78KNG</b> | PGXome Custom - Mitochondrial Complex IV Deficiency  | PreventionGenetics, part of Exact Sciences |
| <b>6P7NG</b> | PGXome Custom - Mitochondrial Complex V Deficiency   | PreventionGenetics, part of Exact Sciences |
| <b>77SZG</b> | Respiratory Chain Deficiency Panel   | Knight Molecular Diagnostic                |
| <b>7B35G</b> | STAT Comprehensive Mitochondrial Nuclear Gene Panel  | PerkinElmer Genomics                       |

| GTU   | Test Name   | Laboratory Name      |
|-------|---|----------------------|
| 7B38G | STAT Mitochondrial Encephalopathy/Leigh Syndrome Nuclear Gene Panel | PerkinElmer Genomics |

### Mitochondrial DNA (mtDNA) Sequencing Panel Tests

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 7SKYG | Advanced mtDNA Point Mutations   | LabCorp   |
| 76WBG | CentoMito® Comprehensive - NGS Panel (Mito Genome included)                          | Centogene   |
| 79ASG | CentoMito® Genome - Mito Genome  | Centogene   |
| 3KDTG | CHOP MitoGenome Sequencing + Deletion Analysis                                       | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 2Z9XG | Complete Mitochondrial Sequencing (mtDNA)  | Gene by Gene  |
| 2DZ4G | Comprehensive mtDNA Analysis by Massively Parallel Sequencing                        | Baylor Genetics, LLC  |
| 79MYG | Deafness, nonsyndromic, sensorineural, mitochondrial - Single Gene Sequencing by NGS | Centogene   |
| 2LKVG | Dual Genome Panel by Massively Parallel Sequencing                                   | Baylor Genetics, LLC  |
| 7V38G | Genomic Unity® Mitochondrial Genome Sequence Analysis                                | Variantyx, Inc.   |
| 39CMG | Known mtDNA Variant(s) Testing by NGS-Urine-Test T822                                | GeneDx  |
| 4AZZG | Maternal MitoGenome Seq + Del  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 6U87G | Maternal Relative Exome + MitoGenome Combined Test                                   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 6L9XG | Mito Genome Sequencing & Deletion Testing  | GeneDx  |
| 7QBEG | Mitochondrial Depletion NGS Panel  | Greenwood Genetic Center - Molecular Diagnostic Laboratory            |
| 253UG | Mitochondrial Disorders (mtDNA) Sequencing and Deletion Analysis by NGS              | ARUP Laboratories   |
| 72G9G | Mitochondrial DNA Depletion Testing (Muscle)   | LabCorp   |
| 5D36G | Mitochondrial DNA Sequencing   | New Jersey Medical School - Institute of Genomic Medicine             |
| 2G4RG | Mitochondrial DNA Testing  | Bionano Laboratories  |
| 6R5TG | Mitochondrial Full Genome Analysis, Next-Generation Sequencing (NGS), Varies         | Mayo Clinic Laboratories  |
| 4CH3G | MITOCHONDRIAL GENOME SEQUENCING  | Center for Human Genetics, Inc.                                       |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 7RXFG | MITOCHONDRIAL GENOME SEQUENCING                                      | LabCorp   |
| 6R9BG | MITOCHONDRIAL GENOME SEQUENCING                                      | MNG Laboratories  |
| 6R9DG | Mitochondrial Genome Sequencing + Deletion Analysis                  | MNG Laboratories  |
| 7RXEG | Mitochondrial Genome Sequencing and Deletion Analysis                | LabCorp   |
| 28H4G | Mitochondrial Myopathy mtDNA   | Johns Hopkins Medical Institutions - Pathology Laboratory             |
| 7WC5G | Mitochondrial Whole Genome Sequence Analysis                         | University of Minnesota Physicians Outreach Laboratory                |
| 2XYNG | mtDNA Whole Genome Sequencing  | Columbia University - Personalized Genomic Medicine                   |
| 27KAG | MVL MitoSeq Panel  | Molecular Vision Laboratory   |
| 7R7XG | Nuclear Mitochondrial Gene Panel, Next-Generation Sequencing, Varies | Mayo Clinic Laboratories  |
| 6USKG | PGmito - Mitochondrial Genome Sequencing                             | PreventionGenetics, part of Exact Sciences                            |
| 35JUG | Rapid MitoGenome Seq + Del   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |

### Pan-Cancer Hereditary Cancer Panel Tests

| GTU   | Test Name  | Laboratory Name                            |
|-------|--|--|
| 7DQMG | BRCA2/CHEK2 Del/Dup Testing by MLPA                        | PerkinElmer Genomics                       |
| 76DVG | Cancer Panel   | PreventionGenetics, part of Exact Sciences |
| 76DGG | CancerNext   | Ambry Genetics                             |
| 6UR6G | CancerNext with RNAinsight                                 | Ambry Genetics                             |
| 7TP4G | CancerNext Expanded HBOC                                   | Ambry Genetics                             |
| 7TP2G | CancerNext: HBOC   | Ambry Genetics                             |
| 76E2G | CancerNext-Expanded  | Ambry Genetics                             |
| 6UR7G | CancerNext-Expanded with RNAinsight                        | Ambry Genetics                             |
| 5BTUG | Centocancer - NGS Panel (CNV included)                     | Centogene                                  |
| 5APDG | Centocancer comprehensive panel - NGS Panel (CNV included) | Centogene                                  |
| 7C5QG | Color Extended   | Color Genomics                             |
| 7C5RG | Color Standard   | Color Genomics                             |



| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 76E5G | Common Cancer Management Panel   | GeneDx  |
| 7V2AG | Common Hereditary Cancer Panel   | Baylor Genetics, LLC  |
| 7RX8G | Common Hereditary Cancer Screening Panel   | PreventionGenetics, part of Exact Sciences                            |
| 76BTG | COMP CANCER PANEL  | ACL Laboratories  |
| 76CKG | Comprehensive Cancer Panel   | PerkinElmer Genomics  |
| 76DQG | Comprehensive Common Cancer Panel  | BioReference Laboratories   |
| 76C3G | Comprehensive Common Cancer Panel: HBOC  | GeneDx  |
| 5FV7G | 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                                  |
| 7V28G | Comprehensive Hereditary Cancer Panel  | Baylor Genetics, LLC  |
| 5K7ZG | Comprehensive Hereditary Cancer Panel  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 262EG | Comprehensive Hereditary Cancer Panel  | Quest Diagnostics   |
| 7BP4G | Comprehensive Hereditary Cancer Panel  | The University of Chicago Genetic Services                            |
| 77HHG | Comprehensive Pan-cancer analysis (DNA and RNA analysis)   | Children's Hospital Colorado Precision Diagnostics Laboratory         |
| 76DHG | CustomNext-Cancer  | Ambry Genetics  |
| 7WJYG | CustomNext-Cancer  | University of Minnesota Physicians Outreach Laboratory                |
| 6UR2G | CustomNext-Cancer with RNAinsight  | Ambry Genetics  |
| 76CAG | Empower - Multi-cancer expanded Hereditary Cancer Test   | Natera  |
| 76C5G | Empower - Multi-cancer Hereditary Cancer Test  | Natera  |
| 4M9NG | Full Comprehensive Cancer Panel (Deletion/Duplication Only)  | Fulgent Genetics  |
| 6U89G | Full Comprehensive Cancer Panel (Germline)   | NeoGenomics Laboratories  |
| 76CUG | Full Comprehensive Cancer Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics  |
| 4U95G | Full Comprehensive Cancer Panel (Sequencing Only)  | Fulgent Genetics  |
| 3HLKG | Full Focus Cancer Panel (Deletion/Duplication Only)  | Fulgent Genetics  |
| 6U8AG | Full Focus Cancer Panel (Germline)   | NeoGenomics Laboratories  |
| 76DMG | Full Focus Cancer Panel (Sequencing & Deletion/Duplication)  | Fulgent Genetics  |

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 4CCHG | Full Focus Cancer Panel (Sequencing Only)   | Fulgent Genetics   |
| 7VZFG | GeneticsNow Comprehensive   | GoPath Laboratories  |
| 25XXG | Guideline Based Hereditary Cancer Panel   | Quest Diagnostics  |
| 7X24G | GxVISION Hereditary Cancer Risk Assessment Comprehensive Inherited Cancer Gene Tests                    | Otogenetics  |
| 6URZG | HC COMMON HEREDITARY CANCER PANEL   | UCSF Molecular Diagnostics Laboratory                                  |
| 76BWG | Hereditary Breast and Ovarian Cancer - Expanded and Lynch Syndrome Panel                                | PreventionGenetics, part of Exact Sciences                             |
| 76DRG | Hereditary Cancer   | Color Genomics   |
| 7VAGG | Hereditary Cancer NGS Panel   | AiLife Diagnostics   |
| 7VAHG | Hereditary Cancer NGS Panel Rapid   | AiLife Diagnostics   |
| 76DXG | Hereditary Cancer Panel, Sequencing and Deletion/Duplication  | ARUP Laboratories  |
| 7DLHG | Hereditary Common Cancer Panel, Varies  | Mayo Clinic Laboratories   |
| 7DKGG | Hereditary Expanded Cancer Panel, Varies  | Mayo Clinic Laboratories   |
| 76CMG | High/mod Risk Panel (20 genes)  | ACL Laboratories   |
| 7VVFG | HopeSeq Germline Confirmatory Panel (Hope Hereditary Cancer Predisposition Panel)                       | City of Hope National Medical Center - Molecular Diagnostic Laboratory |
| 76DKG | Inherited Cancer Panel  | Knight Molecular Diagnostic  |
| 76CHG | Integrated BRACAnalysis with Myriad myRisk Hereditary Cancer Update Test                                | Myriad Genetics  |
| 77BJG | Invitae Cancer Screen   | Invitae Corporation  |
| 7UZRG | Invitae Common Hereditary Cancers + RNA Panel   | Invitae Corporation  |
| 7DGXG | Invitae Common Hereditary Cancers + RNA Panel-Common Hereditary Cancers Genes Eligible for RNA Analysis | Invitae Corporation  |
| 76BNG | Invitae Common Hereditary Cancers Panel   | Invitae Corporation  |
| 7UZSG | Invitae Multi-Cancer + RNA Panel  | Invitae Corporation  |
| 7AV8G | Invitae Multi-Cancer + RNA Panel-Multi-Cancer Genes Eligible for RNA Analysis                           | Invitae Corporation  |
| 76BMG | Invitae Multi-Cancer Panel  | Invitae Corporation  |
| 76BUG | Myriad myRisk   | Myriad Genetics  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 6RMUG | myRisk Update Test  | Myriad Genetics   |
| 7DQGG | NxGen Hereditary Cancer Panel   | NxGen MDx   |
| 76CWG | OncoGene Dx Custom Panel  | GeneDx  |
| 7V3LG | OnkoRisk Hereditary Oncology Guideline Panel (Non-NYS)                                    | BioReference Laboratories                                     |
| 7V3MG | OnkoRisk Hereditary Oncology Management Panel (Non-NYS)                                   | BioReference Laboratories                                     |
| 7V3KG | OnkoRisk Hereditary Oncology Plus Panel (Non-NYS)   | BioReference Laboratories                                     |
| 78FJG | Pan-cancer DNA Analysis   | Children's Hospital Colorado Precision Diagnostics Laboratory |
| 2TQXG | PrevenTest  | Advanced Molecular Diagnostics, LLC                           |
| 74G2G | Rest of Comprehensive Common Cancer Panel: HBOC   | GeneDx  |
| 7RVTG | Riskguard   | Exact Sciences Laboratories, LLC                              |
| 7B7FG | STAT Comprehensive Cancer Panel   | PerkinElmer Genomics  |
| 7TS6G | Targeted Oncology Panel Next Generation Sequencing Bone Marrow                            | Cleveland Clinic Laboratories                                 |
| 7TS7G | Targeted Oncology Panel Next Generation Sequencing Cytology                               | Cleveland Clinic Laboratories                                 |
| 7TS9G | Targeted Oncology Panel Next Generation Sequencing Other                                  | Cleveland Clinic Laboratories                                 |
| 7VPJG | Tempus xG Common Hereditary Cancers 36 genes (all other hereditary cancer indications)    | Tempus AI, Inc.   |
| 7VPFG | Tempus xG+ Extended Hereditary Cancers 88 genes (all other hereditary cancer indications) | Tempus AI, Inc.   |
| 76D8G | VistaSeq Hereditary Cancer Panel  | Integrated Genetics   |
| 76BXG | VistaSeq Hereditary Cancer Panel  | LabCorp   |
| 76BRG | VistaSeq Hereditary Cancer Panel  | Labcorp   Oncology  |

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

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 7T3YG | Comprehensive Pharmacogenetics Panel                                    | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 2FGMG | Drug Metabolizing Enzymes Panel (CYP2D6, CYP2C9, CYP2C19), DNA analysis | Shodair Children's Hospital - Genetics Laboratory  |
| 7VN9G | EffectiveRx Neuropsychiatric Panel                                      | GENETWORx  |

| GTU   | Test Name   | Laboratory Name  |
|-------|---|--|
| 2AUXG | GeneSight Psychotropic                                      | Myriad Genetics  |
| 5YRYG | GeneSight Psychotropic                                      | Myriad Neuroscience  |
| 33Y2G | Genetic Pharmacology Psychiatry Drug Panel                  | Cincinnati Children's Hospital Medical Center - Molecular Genetics and Cytogenetics Laboratories |
| 6S83G | Genomind PGx Test   | Genomind   |
| 7S9DG | IDgenetix   | Castle Biosciences Inc   |
| 7WZVG | Otogenetics PGx Mental Health                               | Otogenetics  |
| 7URYG | PGx: Mental Health Panel                                    | Helix  |
| 7QENG | Pharmacogenetics Panel: Psychotropics                       | ARUP Laboratories  |
| 7UNVG | Pharmacogenetics Panel: Psychotropics, with GeneDose Access | ARUP Laboratories  |
| 7PRTG | Psych HealthPGx Panel                                       | RPRD Diagnostics LLC   |
| 6UAFG | Psychotropic Pharmacogenomics Gene Panel, Varies            | Mayo Clinic Laboratories   |
| 3WG4G | SureGene  | Clinical Reference Laboratory  |
| 33HHG | Tempus  nP  | Tempus AI, Inc.  |

### Prostate Cancer Prognostic Algorithmic Tests

| GTU   | Test Name                            | Laboratory Name |
|-------|--------------------------------------|-----------------|
| 7BN5G | Decipher Prostate Genomic Classifier | Veracyte        |
| 7Q8UG | Genomic Prostate Score® Test         | MDx Health      |
| 6UUUG | Prolaris                             | Myriad Genetics |
| 6UU7G | Prolaris Biopsy                      | Myriad Genetics |

### Rapid Genome Sequencing Tests

| GTU   | Test Name  | Laboratory Name |
|-------|--|-----------------|
| 7VRKG | GenomeXpress® - Duo - Rapid Genome Sequencing with Provisional Results in 7 Days     | GeneDx          |
| 7VRLG | GenomeXpress® - Proband - Rapid Genome Sequencing with Provisional Results in 7 Days | GeneDx          |

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 6UNRG | GenomeExpress® - Trio - Rapid Genome Sequencing with Provisional Results in 7 Days | GeneDx  |
| 6R9TG | MNGenome STAT DUO Sequencing   | MNG Laboratories  |
| 6R9RG | MNGenome STAT Proband Only Sequencing  | MNG Laboratories  |
| 6R9NG | MNGenome STAT TRIO Sequencing  | MNG Laboratories  |
| 3VAEG | PGnome Rapid   | PreventionGenetics, part of Exact Sciences                            |
| 3CSHG | PGnome® - RAPID - Duo  | PreventionGenetics, part of Exact Sciences                            |
| 3VD3G | PGnome® - RAPID - Trio   | PreventionGenetics, part of Exact Sciences                            |
| 7V9ZG | Rapid Duo WGS  | AiLife Diagnostics  |
| 86SYG | Rapid Duo Whole Genome Sequencing  | Baylor Genetics, LLC  |
| 7V9VG | Rapid Proband WGS  | AiLife Diagnostics  |
| 86SZG | Rapid Proband Whole Genome Sequencing  | Baylor Genetics, LLC  |
| 7U3JG | Rapid Targeted Analysis of Family Member   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 7V9XG | Rapid Trio WGS   | AiLife Diagnostics  |
| 7AUWG | Rapid Trio Whole Genome Sequencing   | Baylor Genetics, LLC  |
| 7T4JG | Rapid Whole Genome Sequencing  | ARUP Laboratories   |
| 7RNXG | Rapid Whole Genome Sequencing Duo  | Rady Children's Institute for Genomic Medicine                        |
| 7RNYG | Rapid Whole Genome Sequencing Proband Only   | Rady Children's Institute for Genomic Medicine                        |
| 7RNZG | Rapid Whole Genome Sequencing Trio   | Rady Children's Institute for Genomic Medicine                        |
| 7T4LG | Rapid Whole Genome Sequencing, Familial Control                                    | ARUP Laboratories   |
| 7T4KG | Rapid Whole Genome Sequencing, Familial Control with Report                        | ARUP Laboratories   |
| 7ZR9G | STAT Prenatal Whole Genome Sequencing, DUO   | PerkinElmer Genomics  |
| 7ZR7G | STAT Prenatal Whole Genome Sequencing, Proband ONLY                                | PerkinElmer Genomics  |
| 7ZR8G | STAT Prenatal Whole Genome Sequencing, QUAD  | PerkinElmer Genomics  |
| 7ZR8G | STAT Prenatal Whole Genome Sequencing, TRIO  | PerkinElmer Genomics  |
| 7RSEG | Ultra-rapid Whole Genome Sequencing  | Rady Children's Institute for Genomic Medicine                        |
| 7U24G | Whole Genome Sequencing, Rapid-Duo (Baylor)  | University of Michigan - Michigan Medical Genetics Laboratories       |
| 7U25G | Whole Genome Sequencing, Rapid-Proband (Baylor)                                    | University of Michigan - Michigan Medical Genetics Laboratories       |

## Thyroid Cancer Diagnostic Algorithmic Tests

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

## TPMT and NUDT15 Typing Tests

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

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

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

| GTU   | Test Name  | Laboratory Name  |
|-------|--|--|
| 2YTJG | BCR / ABL - t(9;22) by RT-PCR (Quantitative)                                   | Labcorp   Oncology   |
| 3LU5G | BCR ABL P210 QUANT   | SUNY Upstate Medical University  |
| 7W5EG | BCR ABL1 Major Breakpoint Quant P210   | University of Minnesota Physicians Outreach Laboratory                 |
| 7BQPG | BCR/ABL by FISH  | Allina Health Laboratory   |
| 6ND3G | BCR/ABL D-FISH   | Center for Genetic Testing at Saint Francis                            |
| 3KP8G | BCR/ABL qRT PCR, Major p210 and Minor p190 Mutations, Minimal Residual Disease | Molecular Pathology Laboratory Network                                 |
| 2V82G | BCR/ABL Quantification by RT-PCR   | BioReference Laboratories  |
| 6YXMG | BCR/ABL t(9;22) major (p210) IS Quantitative                                   | Molecular Diagnostic Laboratory - Barnes Jewish Hospital               |
| 6LE9G | BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation                        | University of Iowa Hospitals and Clinics - Department of Pathology     |
| 6UV4G | BCR/ABL1 (T(9;22)) RNA Quantitative with Interpretation                        | University of Iowa Hospitals and Clinics - Department of Pathology     |
| 7VVVG | BCR/ABL1 Kinase Mutation Analysis by NGS                                       | City of Hope National Medical Center - Molecular Diagnostic Laboratory |
| 5SUVG | BCR/ABL1 p190 Quantitative PCR Blood   | Cleveland Clinic Laboratories  |
| 5UA8G | BCR/ABL1 p190 Quantitative PCR Bone Marrow                                     | Cleveland Clinic Laboratories  |
| 7TH5G | BCR/ABL1 p210 and p190 Diagnostic PCR Blood                                    | Cleveland Clinic Laboratories  |
| 7TH6G | BCR/ABL1 p210 and p190 Diagnostic PCR Bone Marrow                              | Cleveland Clinic Laboratories  |
| 5UAAG | BCR/ABL1 p210 Quantitative PCR Blood   | Cleveland Clinic Laboratories  |
| 6LQRG | BCR/ABL1 p210 Quantitative PCR Bone Marrow                                     | Cleveland Clinic Laboratories  |
| 5PH3G | BCR-ABL Diagnostic, Blood  | Johns Hopkins Medical Institutions - Pathology Laboratory              |
| 4GZ7G | BCR-ABL Diagnostic, Bone Marrow  | Johns Hopkins Medical Institutions - Pathology Laboratory              |
| 5W36G | BCR-ABL Mutation   | University of Illinois at Chicago - Biochemical Genetics Laboratory    |
| 5PGYG | BCR-ABL p190 Followup, Blood   | Johns Hopkins Medical Institutions - Pathology Laboratory              |
| 4GLHG | BCR-ABL p190 Followup, Bone Marrow   | Johns Hopkins Medical Institutions - Pathology Laboratory              |
| 4AGMG | BCR-ABL p210 Followup, Blood   | Johns Hopkins Medical Institutions - Pathology Laboratory              |

| GTU    | Test Name   | Laboratory Name  |
|--------|---|--|
| 3AUMG  | BCR-ABL p210 Followup, Bone Marrow  | Johns Hopkins Medical Institutions - Pathology Laboratory                                    |
| 723HG  | BCR-ABL1 Gene Rearrangement, Quantitative, PCR  | Quest Diagnostics  |
| 7WVTG  | BCR-ABL1 High Sensitivity Major p210  | University of Minnesota Physicians Outreach Laboratory                                       |
| 8E7CG  | BCR-ABL1 Major (p210) by Quantitative RT-PCR for Monitoring   | Medfusion  |
| 8E7GG  | BCR-ABL1 Major (p210) by Quantitative RT-PCR with Reflex to Minor (p190) for Baseline Quantitation                      | Medfusion  |
| 2ZTAG  | BCR-ABL1 Non-Standard p230  | NeoGenomics Laboratories   |
| 7BD CG | BCR-ABL1 QUAL, RFLX QUANT, NEW DIAGNOSIS  | Clinical Pathology Laboratories  |
| 7AEXG  | BCR-ABL1 QUAL, RFLX QUANT, NEW DIAGNOSIS, WHOLE BLOOD   | Clinical Pathology Laboratories  |
| 7AF2G  | BCR-ABL1 QUANT, MAJOR, WHOLE BLOOD  | Clinical Pathology Laboratories  |
| 7AEZG  | BCR-ABL1 QUANT, MINOR, WHOLE BLOOD  | Clinical Pathology Laboratories  |
| 73BFG  | BCR-ABL1 Standard p210, p190  | NeoGenomics Laboratories   |
| 6X7DG  | BCR-ABL1 Transcript Detection for Chronic Myelogenous Leukemia (CML) and Acute Lymphocytic Leukemia (ALL), Quantitative | LabCorp  |
| 86NEG  | BCR-ABL1, Major (p210), Quantitative  | Molecular Diagnosis - University of Rochester Medical Center - Strong Memorial Hospital      |
| 7BDEG  | BCR-ABL1, QUANT, CML MONITORING (p210)  | Clinical Pathology Laboratories  |
| 7BDAG  | BCR-ABL1, QUANT, MINOR (p190)   | Clinical Pathology Laboratories  |
| 5YZ4G  | FISH (9_22)   | SUNY Upstate Medical University  |
| 6VXDG  | QUANTITATIVE DETECTION OF BCR-ABL1 MAJOR FORM (p210)  | University of California Davis Health System - UCDCMC - Molecular and Cytogenetic Laboratory |

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

| GTU   | Test Name  | Laboratory Name   |
|-------|--|---|
| 86SBG | Altera Tumor Profiling                           | Natera  |
| 3CQFG | Columbia Combined Cancer Panel (CCCP)            | Columbia University - Personalized Genomic Medicine           |
| 2CV6G | Comprehensive NGS Solid Tumor Mutation Panel     | UCSD - Molecular Diagnostics & Cytogenetics Laboratory        |
| 2D5CG | Comprehensive Solid Tumor (DNA and RNA analysis) | Children's Hospital Colorado Precision Diagnostics Laboratory |



| <b>GTU</b>   | <b>Test Name</b>  | <b>Laboratory Name</b>  |
|--------------|---|---|
| <b>7VQWG</b> | Comprehensive Solid Tumor Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| <b>72Y9G</b> | Endeavor Comprehensive Solid Tumor Profile                                | PathGroup   |
| <b>6R4AG</b> | FoundationOne CDx   | Foundation Medicine   |
| <b>7UJGG</b> | Fusion Transcript Panel   | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory    |
| <b>3V7ZG</b> | GatorSeq (NGS Panel)  | University of Florida - Health Pathology Laboratories                             |
| <b>7XMNG</b> | GeneAssure Solid Tumor NGS Assay  | University of Illinois at Chicago - Biochemical Genetics Laboratory               |
| <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>86PJG</b> | Guardant360 TissueNext  | Guardant Health   |
| <b>6ZQMG</b> | Iowa Cancer Mutation and RNA Fusion Profile and Interpretation            | University of Iowa Hospitals and Clinics - Department of Pathology                |
| <b>7AWEG</b> | MayoComplete Solid Tumor Panel, Next-Generation Sequencing, Tumor         | Mayo Clinic Laboratories  |
| <b>6WUTG</b> | MI Tumor Seek Hybrid  | Caris Life Sciences   |
| <b>6UUMG</b> | MSK-IMPACT  | Memorial Sloan Kettering Cancer Center  |
| <b>7SSDG</b> | Neo Comprehensive - Solid Tumor   | NeoGenomics Laboratories  |
| <b>7529G</b> | NeXT Dx™  | Personalis  |
| <b>34WBG</b> | NGS Solid Tumor Panel   | Johns Hopkins Medical Institutions - Pathology Laboratory                         |
| <b>7V34G</b> | OncoAlly Solid Tumor Analysis   | Varietyx, Inc.  |
| <b>7BNDG</b> | OncoExTra®  | Exact Sciences Laboratories, LLC  |
| <b>2YA6G</b> | OncoGxOne   | Admera Health   |
| <b>76ETG</b> | Oncoplex Select Cancer Gene Panel   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>76EQG</b> | Oncoplex Select Director  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>76EYG</b> | Oncoplex Select Interpretation  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>76EZG</b> | Oncoplex Select Methods   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |

| <b>GTU</b>   | <b>Test Name</b>  | <b>Laboratory Name</b>  |
|--------------|---|---|
| <b>76ESG</b> | Oncoplex Select Panel   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>76ERG</b> | Oncoplex Select Result  | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>76EXG</b> | Oncoplex Select Tested Sample   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| <b>7UUYG</b> | OnkoSight Advanced 523 Gene NGS Panel   | BioReference Laboratories   |
| <b>7SSVG</b> | OnkoSight Advanced 523 Gene NGS Reanalysis with PierianDx                         | BioReference Laboratories   |
| <b>7URPG</b> | OnkoSight Advanced 523 Gene NGS with PierianDx Interpretation                     | BioReference Laboratories   |
| <b>2FLHG</b> | Paired Tumor/Normal – Comprehensive Solid Tumor Panel                             | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| <b>2FLGG</b> | Paired Tumor/Normal – Solid Tumor Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| <b>7TKLG</b> | Pan-Cancer Solid Tumor NGS Panel  | UCLA Diagnostic Molecular Pathology Laboratory                                    |
| <b>6V93G</b> | PCMP - Personalized Cancer Mutation Panel   | University of Pittsburgh Medical Center - Division of Molecular Diagnostics       |
| <b>7UJFG</b> | PennSeq Solid Tumor Panel   | University of Pennsylvania School of Medicine - Molecular Pathology Laboratory    |
| <b>2K9WG</b> | PGDx elio tissue complete   | Personal Genome Diagnostics   |
| <b>7A5HG</b> | Precise Tumor   | Myriad Genetics   |
| <b>2DSFG</b> | RNA Fusion Analysis   | Children's Hospital Colorado Precision Diagnostics Laboratory                     |
| <b>7ATMG</b> | SOLID TUMOR COMPREHENSIVE PANEL (Combined solid tumor mutation and fusion panels) | Texas Children's Hospital   |
| <b>2D5BG</b> | Solid Tumor DNA Analysis  | Children's Hospital Colorado Precision Diagnostics Laboratory                     |
| <b>7SVMG</b> | Solid Tumor Expanded Panel  | Quest Diagnostics   |
| <b>3CSSG</b> | Solid Tumor Molecular Profile   | Fulgent Genetics  |
| <b>2NSPG</b> | Solid Tumor NGS Panel   | University of Michigan - Michigan Medical Genetics Laboratories                   |
| <b>2FRPG</b> | Solid Tumor Panel   | Children's Hospital of Philadelphia - Division of Genomic Diagnostics             |
| <b>6UWNG</b> | Solid tumor panel - Somatic Mutation Analysis                                     | Centogene   |
| <b>2LEVG</b> | Solid Tumor Targeted Cancer Gene Panel by Next-Generation Sequencing, NGS         | Medical College of Wisconsin - Human and Molecular Genetics Center                |
| <b>7C3KG</b> | Solid tumor targeted NGS panel  | Allina Health Laboratory  |

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 7UQMG | Solid TumorSEQ Expanded Panel                             | Medfusion   |
| 2CPUG | Strata Select   | Strata Oncology   |
| 7TXFG | Tempus xT (tumor only), Paraffin Block                    | University of Michigan - Michigan Medical Genetics Laboratories                   |
| 7TXHG | Tempus xT Targeted Panel (Tumor plus Blood)               | University of Michigan - Michigan Medical Genetics Laboratories                   |
| 7TXJG | Tempus xT Targeted Panel (Tumor plus Saliva)              | University of Michigan - Michigan Medical Genetics Laboratories                   |
| 7VPEG | Tempus xT CDx: FDA-approved Companion Diagnostic          | Tempus AI, Inc.   |
| 86SDG | Tempus xT: Targeted panel of 648 genes (Solid Tumor Only) | Tempus AI, Inc.   |
| 6RWJG | TempusTM Test   | ACL Laboratories  |
| 6LS8G | UW OncoPlex Cancer Gene Panel                             | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |
| 5LYGG | UW OncoPlex Single Gene                                   | University of Washington Medical Center - Laboratory Medicine-Genetics Laboratory |

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

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 2LVRG | Cancer Mutation Profiling and Interpretation          | University of Iowa Hospitals and Clinics - Department of Pathology          |
| 2LCCG | Cancer Mutation Profiling Blood Paired Normal Testing | University of Iowa Hospitals and Clinics - Department of Pathology          |
| 2XYRG | Columbia Solid Tumor (CSTP) Subpanels                 | Columbia University - Personalized Genomic Medicine                         |
| 5BZJG | Columbia Solid Tumor Panel (CSTP)                     | Columbia University - Personalized Genomic Medicine                         |
| 7BZJG | GIST targeted NGS panel                               | Allina Health Laboratory  |
| 5BRWG | NGS Solid Tumor Hotspot Panel                         | Johns Hopkins Medical Institutions - Pathology Laboratory                   |
| 2V3FG | OncoSeq - NGS Panel for Solid Tumors                  | University of Pittsburgh Medical Center - Division of Molecular Diagnostics |
| 86LTG | OnkoSight Advanced Solid Tumor NGS Panel              | BioReference Laboratories   |
| 4ZPLG | Solid Tumor 15 Genes Panel by NGS                     | Beaumont Laboratories - Molecular Pathology Lab                             |
| 2E5TG | Solid Tumor Core Panel                                | Quest Diagnostics   |
| 6VAPG | Solid Tumor Mutation Panel                            | Ohio State University - Molecular Pathology Laboratory                      |
| 6V3PG | Solid Tumor Mutation Panel 15                         | ACL Laboratories  |
| 7DK3G | Solid Tumor Mutation Panel, Sequencing                | ARUP Laboratories   |

| GTU   | Test Name                             | Laboratory Name   |
|-------|---------------------------------------|---|
| 7PVTG | Solid Tumor Precision Panel           | Nebraska Medical Center - Molecular Diagnostic Laboratory |
| 7UQNG | Solid TumorSEQ Core Panel             | Medfusion   |
| 7XWDG | Tumor (NOS) - Custom Panel - 5 Probes | John Hopkins All Children's Hospital                      |

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

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

### Uveal Melanoma Prognostic Algorithmic Tests

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

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

| GTU   | Test Name   | Laboratory Name   |
|-------|---|---|
| 2FM3G | 90-gene XLID Panel  | Children's Hospital of Philadelphia - Division of Genomic Diagnostics |
| 7V35G | Genomic Unity® X-linked Intellectual Disability Plus Analysis | Varietyx, Inc.  |
| 4XMCG | PANEL 2 (ATPGAP2, OPHN1, SLC6A8, SYN1) SEQUENCING ONLY        | Center for Human Genetics, Inc.                                       |
| 4SMHG | PANEL 2 MLPA (OPHN1, SLC6A8)                                  | Center for Human Genetics, Inc.                                       |

| <b>GTU</b>   | <b>Test Name</b>   | <b>Laboratory Name</b>                                     |
|--------------|--|--|
| <b>78KLG</b> | PGXome Custom - Intellectual Disability, X-linked  | PreventionGenetics, part of Exact Sciences                 |
| <b>23XZG</b> | XLID (X-Linked Intellectual Disability) NGS Panel (Deletion/Duplication Only)                                    | Fulgent Genetics   |
| <b>5DSSG</b> | XLID (X-Linked Intellectual Disability) NGS Panel (Sequencing & Deletion/Duplication)                            | Fulgent Genetics   |
| <b>34N9G</b> | XLID (X-Linked Intellectual Disability) NGS Panel (Sequencing Only)  | Fulgent Genetics   |
| <b>4S99G</b> | XLID TIER 1 SEQUENCING   | Center for Human Genetics, Inc.                            |
| <b>3KWVG</b> | XLID TIER 2 MLPA   | Center for Human Genetics, Inc.                            |
| <b>3KMHG</b> | XLID TIER 2 SEQUENCING   | Center for Human Genetics, Inc.                            |
| <b>5YXXG</b> | XLID TIER 3 MLPA ONLY  | Center for Human Genetics, Inc.                            |
| <b>2BSYG</b> | XLID TIER 3 SEQUENCING ONLY  | Center for Human Genetics, Inc.                            |
| <b>5XPVG</b> | XLID TIER 4 MLPA   | Center for Human Genetics, Inc.                            |
| <b>55YDG</b> | XLID TIER 4 SEQUENCING ONLY  | Center for Human Genetics, Inc.                            |
| <b>2BYZG</b> | X-linked Intellectual Disability (NGS Panel and Copy Number Analysis + Fragile X Repeat Expansion & Methylation) | MNG Laboratories   |
| <b>33HUG</b> | X-Linked Intellectual Disability (XLID) NGS Panel  | Greenwood Genetic Center - Molecular Diagnostic Laboratory |
| <b>4YJGG</b> | X-linked Intellectual Disability Panel   | PreventionGenetics, part of Exact Sciences                 |
| <b>7727G</b> | X-Linked Non-Specific Intellectual Disability Panel  | The University of Chicago Genetic Services                 |

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