UNICARE STATE INDEMNITY PLAN

PREAPPROVALS – GENETIC TESTING

For UniCare State Indemnity Plan/Basic, PLUS and Community Choice members

Effective July 1, 2019

For preapprovals

AIM Specialty Health
866-766-0247
www.providerportal.com

Genetic testing that needs preapproval

- Analysis of KRAS Status
- Analysis of PIK3CA Status in Tumor Cells
- BCR-ABL Mutation Analysis
- BRAF Mutation Analysis
- Cardiac Ion Channel Genetic Testing
- Cell-Free Fetal DNA-Based Prenatal Screening for Fetal Aneuploidy
- Chromosomal Microarray Analysis (CMA) for Developmental Delay, Autism Spectrum Disorder, Intellectual Disability (Intellectual Developmental Disorder) and Congenital Anomalies
- Detection and Quantification of Tumor DNA Using Next Generation Sequencing in Lymphoid Cancers
- DNA-Based Testing for Adolescent Idiopathic Scoliosis
- Epidermal Growth Factor Receptor (EGFR) Testing
- Gene-Based Tests for Screening, Detection and Management of Prostate Cancer
- Gene Expression Profile Tests for Multiple Myeloma
- Gene Expression Profiling for Cancers of Unknown Primary Site
- Gene Expression Profiling for Colorectal Cancer
- Gene Expression Profiling for Managing Breast Cancer Treatment
- Gene Expression Profiling of Melanomas
- Genetic Testing and Biochemical Markers for the Diagnosis of Alzheimer's Disease
- Genetic Testing for Breast and/or Ovarian Cancer Syndrome
- Genetic Testing for Cancer Susceptibility
- Genetic Testing for Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy (CADASIL) Syndrome
- Genetic Testing for CHARGE Syndrome

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Genetic testing that needs preapproval *(continued)*

- Genetic Testing for Colorectal Cancer Susceptibility
- Genetic Testing for Diagnosis and Management of Hereditary Cardiomyopathies (including ARVD/C)
- Genetic Testing for Endocrine Gland Cancer Susceptibility
- Genetic Testing for Frontotemporal Dementia (FTD)
- Genetic Testing for Hereditary Pancreatitis
- Genetic Testing for Inherited Peripheral Neuropathies
- Genetic Testing for Macular Degeneration
- Genetic Testing for PTEN Hamartoma Tumor Syndrome
- Genetic Testing for Statin-Induced Myopathy
- Genetic Testing for TP53 Mutations (Li-Fraumeni Syndrome)
- Genetic Testing of an Individual's Genome for Inherited Diseases
- Genotype Testing for Genetic Polymorphisms to Determine Drug-Metabolizer Status
- In Vitro Companion Diagnostic Devices
- Janus Kinase 2 (JAK2) V617F Gene Mutation Assay
- Methylenetetrahydrofolate Reductase Mutation Testing
- Molecular Marker Evaluation of Thyroid Nodules
- Molecular Profiling for the Evaluation of Malignant Tumors
- The Panexia™ Test for Oncologic Indications
- Preconceptional or Prenatal Genetic Testing of a Parent or Prospective Parent
- Preimplantation Genetic Diagnosis Testing
- Prothrombin G20210A (Factor II) Mutation Testing
- SensiGene® Fetal RhD Genotyping Test
- Short Tandem Repeat Analysis for Specimen Provenance Testing