

Beneficiary Full Name: _____

Sponsor's SSN: _____ - _____ - _____

Date of Birth: _____

Beneficiary State of Residence: _____

Dear Provider,

Please complete and sign this **Single Gene Laboratory Developed Test (LDT) Letter of Attestation** and return as indicated on the additional information request letter or attach it to your online request. TRICARE Operations Manual, Chapter 18, allows coverage for LDTs when specific coverage criteria are met.

This Letter of Attestation is for single-gene LDTs. If you are requesting a panel LDT, please complete and submit the **Panel LDT Letter of Attestation**.

SECTION I – Laboratory developed tests that may be considered for coverage under the Defense Health Agency (DHA) Evaluation of Non-United States Food and Drug Administration (FDA) Approved LDT Demonstration Project. (If the requested test is not indicated in Section I, please complete Section II.)

DIRECTIONS: Mark the single gene(s) test(s) being requested below in COLUMN I, mark the indication for the test in COLUMN II, and write the CPT® code(s) and quantity in COLUMN III being requested.

Failure to complete the form in its entirety will result in a delay in processing your request.

COLUMN I Select the gene(s) being requested	COLUMN II Select the indication(s) for the requested test	COLUMN III Indicate the CPT® code(s) and quantity	
		CPT® code(s)	QTY
<input type="checkbox"/> Afirma Thyroid FNA Analysis	<input type="checkbox"/> To aid in thyroid nodule diagnosis by reducing unnecessary surgeries in patients with indeterminate thyroid nodules. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ALK	<input type="checkbox"/> To determine response to Tyrosine Kinase Inhibitor (TKI) therapy in patients with adenocarcinoma of the lung or mixed lung cancer with adenocarcinoma component of the lung. <input type="checkbox"/> Other indication		
<input type="checkbox"/> APC	<input type="checkbox"/> Testing for APC variants in individuals with clinical symptoms consistent with Familial Adenomatous Polyposis (FAP). <input type="checkbox"/> Testing for APC variants in individuals with clinical symptoms consistent with Attenuated Familial Adenomatous Polyposis (AFAP). <input type="checkbox"/> Testing for APC variants in individuals with clinical symptoms consistent with Turcot's or Gardner's syndromes. <input type="checkbox"/> Testing individuals with an APC-associated polyposis syndrome for the purpose of identifying a variant that may be used to screen at-risk relatives. <input type="checkbox"/> For the presymptomatic testing of at-risk relatives for a known familial variant. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ATXN1	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 1 (SCA1) in patients with cerebellar ataxia of unknown etiology, along with extracerebellar symptoms associated with SCA1 and/or a family history consistent with autosomal dominant inheritance. <input type="checkbox"/> Diagnosis of SCA1 in symptomatic family members of known SCA1 patients. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> ATXN2	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 2 (SCA2) in patients with cerebellar ataxia of unknown etiology, along with extracerebellar symptoms associated with SCA2 and/or a family history consistent with autosomal dominant inheritance. <input type="checkbox"/> Diagnosis of SCA2 in symptomatic family members of known SCA2 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ATXN3	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 3 (SCA3) in patients with cerebellar ataxia of unknown etiology, along with extracerebellar symptoms associated with SCA3 and/or a family history consistent with autosomal dominant inheritance. <input type="checkbox"/> Diagnosis of SCA3 in symptomatic family members of known SCA3 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ATXN7	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 7 (SCA7) in patients with cerebellar ataxia and visual disturbance. <input type="checkbox"/> Diagnosis of SCA7 in symptomatic family members of known SCA7 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ATXN10	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 10 (SCA10) in ataxia patients whose ancestry is of American Indian origin, and whose family history is consistent with autosomal dominant inheritance. <input type="checkbox"/> Diagnosis of SCA10 in symptomatic family members of known SCA10 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> BCR/ABL1	<input type="checkbox"/> Diagnostic assessment of individuals with suspected Chronic Myelogenous Leukemia (CML) by quantitative RT-PCR (RQ-PCR). <input type="checkbox"/> Diagnostic assessment of individuals with suspected CML by qualitative RT-PCR. <input type="checkbox"/> Monitoring response to TKI therapy, such as imatinib, in individuals with CML by RQ-PCR. <input type="checkbox"/> Testing for the presence of the BCR/ABL1 p.Thr315Ile variant in CML patients to guide treatment selection following resistance to first-line imatinib therapy. <input type="checkbox"/> Testing for the presence of BCR/ABL1 variants other than p.Thr315Ile in CML patients to guide treatment selection following resistance to first-line imatinib therapy. <input type="checkbox"/> Other indication		
<input type="checkbox"/> BMPR1A	<input type="checkbox"/> To clarify the diagnosis of individuals with Juvenile Polyposis Syndrome (JPS). <input type="checkbox"/> If a known SMAD4 mutation is in the family, genetic testing should be performed in the first six months of life due to hereditary hemorrhagic telangiectasia risk. <input type="checkbox"/> Other indication		
<input type="checkbox"/> BRAF	<input type="checkbox"/> To predict response to vemurafenib therapy in patients with a positive cobas 4800 BRAF mutation test result. <input type="checkbox"/> To predict response to trametinib monotherapy in advanced melanoma patients with a positive BRAF p.Val600Glu and/or p.Val600Lys test result. <input type="checkbox"/> To predict response to dabrafenib monotherapy in advanced melanoma patients with a positive BRAF p.Val600Glu test result. <input type="checkbox"/> To predict response to trametinib and dabrafenib combination therapy in advanced melanoma patients with a positive BRAF p.Val600Glu and/or p.Val600Lys test result. <input type="checkbox"/> For individuals with indeterminate thyroid Fine-Needle Aspiration (FNA) biopsy cytology for diagnosis of papillary thyroid carcinoma. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> BRCA1/BRCA2	Please provide the clinical indication(s) for BRCA1/BRCA2 gene testing:		
<input type="checkbox"/> CACNA1A	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 6 (SCA6) in patients with cerebellar ataxia with dysarthria and/or nystagmus. <input type="checkbox"/> Diagnosis of SCA6 in symptomatic family members of known SCA6 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> CALM1, CASQ2, RYR2, and/or TRDN	<input type="checkbox"/> To confirm a diagnosis of Catecholaminergic Polymorphic Ventricular Tachycardia (CPVT) in patients with clinically diagnosed or suspected CPVT. <input type="checkbox"/> Other indication		
<input type="checkbox"/> CDH1	<input type="checkbox"/> For large rearrangements in the CDH1 gene for the treatment of Hereditary Diffuse Gastric Cancer (HDGC). <input type="checkbox"/> Other indication		
<input type="checkbox"/> CEBPA	<input type="checkbox"/> To guide the treatment decisions for individuals with Acute Myeloid Leukemia (AML). <input type="checkbox"/> Other indication		
<input type="checkbox"/> CFTR	<input type="checkbox"/> Confirmation of diagnosis in individuals showing clinical symptoms of Cystic Fibrosis (CF) or having a high sweat chloride level. <input type="checkbox"/> Identification of newborns who are affected with CF. <input type="checkbox"/> Identification of individuals with the p.Gly551Asp variant who will respond to treatment with ivacaftor. <input type="checkbox"/> Male infertility testing and treatment. <input type="checkbox"/> Preconception and prenatal carrier screening in accordance with the most current ACOG guidelines. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Chimerism Analysis	<input type="checkbox"/> For the management and treatment of stem cell transplant patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Chromosome 22q11.2	<input type="checkbox"/> Confirmation of diagnosis in an individual suspected of chromosome 22q11.2 deletion syndrome based on clinical findings. <input type="checkbox"/> Other indication		
<input type="checkbox"/> COL1A1/COL1A2	<input type="checkbox"/> For sequence variants in the COL1A1/COL1A2 genes for the diagnosis of Osteogenesis Imperfecta (OI) when clinical and radiological examination and family history provide inadequate information for diagnosis of OI. <input type="checkbox"/> Other indication		
<input type="checkbox"/> COL3A1	<input type="checkbox"/> To confirm or establish a diagnosis of Ehlers-Danlos Syndrome Type 4 (EDS IV), also known as vascular EDS, in patients with clinical symptoms or features of EDS IV. <input type="checkbox"/> Other indication		
<input type="checkbox"/> CYP2C9	<input type="checkbox"/> For the initiation and management of warfarin treatment. <input type="checkbox"/> Other indication		
<input type="checkbox"/> CYP2C19	<input type="checkbox"/> To manage dosing of clopidogrel. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Cytogenomic Constitutional Microarray Analysis	<input type="checkbox"/> Diagnostic evaluation of patients suspected of having a genetic syndrome (i.e., have congenital anomalies, dysmorphic features, Developmental Delay (DD), and/or intellectual disability). <input type="checkbox"/> Diagnostic evaluation of individuals with Autism Spectrum Disorder (ASD), including autism, Asperger syndrome, and pervasive developmental disorder. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> DAZ/SRY	<input type="checkbox"/> To detect submicroscopic deletions involving the Y chromosome in the evaluation of men with infertility secondary to azoospermia, oligozoospermia, or teratozoospermia. <input type="checkbox"/> Other indication		
<input type="checkbox"/> DMD	<input type="checkbox"/> For diagnostic DMD testing (deletion and duplication analysis with reflex to complete gene sequencing) in males or females exhibiting symptoms of Duchenne Muscular Dystrophy (DMD) or Becker Muscular Dystrophy (BMD). <input type="checkbox"/> Other indication		
<input type="checkbox"/> DMPK	<input type="checkbox"/> Confirmation of a diagnosis of Myotonic Dystrophy Type 1 (DM1) or Type 2 (DM2) in symptomatic patients. <input type="checkbox"/> Diagnosis of DM1 or DM2 in asymptomatic adults who are at an increased risk of DM1 or DM2 through a positive family history. <input type="checkbox"/> Other indication		
<input type="checkbox"/> DSC2, DSG2, DSP, JUP, PKP2, RYR2, TGFB3, and/or TMEM43	<input type="checkbox"/> For sequence variants in the DSC2, DSG2, DSP, JUP, PKP2, RYR2, TGFB3, and TMEM43 genes to confirm a diagnosis of Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C) in probands. <input type="checkbox"/> For a known familial sequence variant in the DSC2, DSG2, DSP, PKP2, or TMEM43 gene for at-risk relatives of probands with International Task Force (ITF)-confirmed ARVD/C to confirm a diagnosis of ARVD/C in those whose symptoms meet the ITF-diagnostic criteria. <input type="checkbox"/> Other indication		
<input type="checkbox"/> DYT1/TOR1A	<input type="checkbox"/> For genetic testing for sequence variants of DYT1 for patients with primary dystonia with onset < 30 years of age. <input type="checkbox"/> For genetic testing for sequence variants of DYT1 for patients with primary dystonia with onset ≥ 30 years of age who have a relative who developed dystonia aged < 30 years. <input type="checkbox"/> Other indication		
<input type="checkbox"/> EGFR	<input type="checkbox"/> To help guide administration of Epidermal Growth Factor Receptor (EGFR) TKIs in the first-line treatment of non-small cell lung cancer. <input type="checkbox"/> Other indication		
<input type="checkbox"/> F2	<input type="checkbox"/> Diagnostic evaluation of individuals with a prior Venous Thromboembolism (VTE) during pregnancy or puerperium. <input type="checkbox"/> For patients with VTE with a personal or family history of recurrent VTE (more than two in the same person). <input type="checkbox"/> For patients with their first VTE before age 50 with no precipitating factors. <input type="checkbox"/> For venous thrombosis at unusual sites such as the cerebral, mesenteric, portal, or hepatic veins. <input type="checkbox"/> For VTE associated with the use of estrogen-containing oral contraceptives, Selective Estrogen Receptor Modulators (SERMs), or Hormone Replacement Therapy (HRT). <input type="checkbox"/> To diagnose an inherited thrombophilia in female family members of individuals with an inherited thrombophilia if the female family member is pregnant or considering pregnancy or oral contraceptive use. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> F5	<input type="checkbox"/> Diagnostic evaluation of individuals with a prior VTE during pregnancy or puerperium. <input type="checkbox"/> For patients with VTE with a personal or family history of recurrent VTE (more than two in the same person). <input type="checkbox"/> For patients with their first VTE before age 50 with no precipitating factors. <input type="checkbox"/> For venous thrombosis at unusual sites such as the cerebral, mesenteric, portal, or hepatic veins. <input type="checkbox"/> For VTE associated with the use of estrogen-containing oral contraceptives, Selective Estrogen Receptor Modulators (SERMs), or Hormone Replacement Therapy (HRT). <input type="checkbox"/> To diagnose an inherited thrombophilia in female family members of individuals with an inherited thrombophilia if the female family member is pregnant or considering pregnancy or oral contraceptive use. <input type="checkbox"/> Other indication		
<input type="checkbox"/> FBN1	<input type="checkbox"/> To facilitate the diagnosis of Marfan syndrome in patients who do not fulfill the Ghent diagnostic criteria, but have at least one major feature of the condition. <input type="checkbox"/> To facilitate the diagnosis of Marfan syndrome in the at-risk relatives of patients carrying known disease-causing variants. <input type="checkbox"/> Other indication		
<input type="checkbox"/> FLCN	<input type="checkbox"/> To confirm a diagnosis of Birt-Hogg-Dubé Syndrome (BHD) in patients with suspected BHD. <input type="checkbox"/> Other indication		
<input type="checkbox"/> FLT3	<input type="checkbox"/> For diagnosis and prognosis in AML. <input type="checkbox"/> Other indication		
<input type="checkbox"/> FMR1	<p>FMR1 gene testing:</p> <input type="checkbox"/> Testing for CGG repeat length for diagnosis of patients of either sex with mental retardation, intellectual disability, developmental delay, or autism. <input type="checkbox"/> Other indication		
	<p>FMR1 gene testing for Fragile X-Associated Tremor/Ataxia Syndrome:</p> <input type="checkbox"/> Males and females older than age 50 years who have progressive cerebellar ataxia and intention tremor with or without a positive family history of FMR1-related disorders in whom other common causes of ataxia have been excluded. <input type="checkbox"/> Women with unexplained Premature Ovarian Insufficiency (POI). <input type="checkbox"/> Other indication		
<input type="checkbox"/> GCK	<input type="checkbox"/> Diagnosis of Maturity-Onset Diabetes of the Young Type 2 (MODY2) in patients with hyperglycemia or non-insulin-dependent diabetes who have a family history of abnormal glucose metabolism in at least two consecutive generations, with the patient or ≥ 1 family member(s) diagnosed before age 25. <input type="checkbox"/> Other indication		
<input type="checkbox"/> GJB2	<input type="checkbox"/> Diagnosis of DFNB1 or DFNA3 in individuals with nonsyndromic hearing loss to aid in treatment. <input type="checkbox"/> Other indication		
<input type="checkbox"/> GJB6	<input type="checkbox"/> Diagnosis of DFNB1 or DFNA3 in individuals with nonsyndromic hearing loss to aid in treatment. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HBA1/HBA2	<input type="checkbox"/> To confirm the diagnosis of alpha-thalassemia in a symptomatic individual. <input type="checkbox"/> To confirm the diagnosis in a pregnant woman with low hemoglobin when alpha-thalassemia is suspected. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> HEXA	<input type="checkbox"/> As an adjunct to biochemical testing in patients with low hexosaminidase A levels in blood. When individuals are identified with apparent deficiency of hexosaminidase A enzymatic activity, targeted mutation analysis can then be used to distinguish pseudodeficiency alleles from disease-causing alleles. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HFE	<input type="checkbox"/> Diagnosis of patients with or without symptoms of iron overload with a serum transferrin saturation >45% and/or elevated serum ferritin. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HLA	<input type="checkbox"/> To determine histocompatibility of tissue between organ and bone marrow donors and recipients prior to transplant. <input type="checkbox"/> For platelet transfusion for patients refractory to treatment due to alloimmunization. <input type="checkbox"/> Diagnosis of celiac disease in symptomatic patients with equivocal results on small bowel biopsy and serology, or in previously symptomatic patients who are asymptomatic while on a gluten-free diet. <input type="checkbox"/> Testing for the HLA-B*1502 allele prior to initiating treatment with carbamazepine in patients from high-risk ethnic groups. <input type="checkbox"/> Testing for the HLA-B*5701 allele for hypersensitivity reactions in patients prior to initiation or reinitiation with treatments containing abacavir. <input type="checkbox"/> Testing for the HLA-B*58:01 allele in patients prior to initiating treatment with allopurinol. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HNF1A	<input type="checkbox"/> Diagnosis of Maturity-Onset Diabetes of the Young Type 3 (MODY3) in patients with hyperglycemia or non-insulin-dependent diabetes who have a family history of abnormal glucose metabolism in at least two consecutive generations, with the patient or ≥ 1 family member(s) diagnosed before age 25. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HNF1B	<input type="checkbox"/> Diagnosis of Maturity-Onset Diabetes of the Young Type 5 (MODY5) in patients with hyperglycemia or non-insulin-dependent diabetes who have a family history of abnormal glucose metabolism in at least two consecutive generations, with the patient or ≥ 1 family member(s) diagnosed before age 25, and who have structural or functional abnormalities of the kidneys. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HNF4A	<input type="checkbox"/> Diagnosis of Maturity-Onset Diabetes of the Young Type 1 (MODY1) in patients with hyperglycemia or non-insulin-dependent diabetes who have a family history of abnormal glucose metabolism in at least two consecutive generations, with the patient or ≥ 1 family member(s) diagnosed before age 25. <input type="checkbox"/> Other indication		
<input type="checkbox"/> HTT	<input type="checkbox"/> To test for CAG repeat length for diagnosis of Huntington Chorea/ Disease (HD) in patients suspected of having HD in the absence of a family history of HD. <input type="checkbox"/> Other indication		
<input type="checkbox"/> IGH	<input type="checkbox"/> For medical management of patients with Acute Lymphoblastic Leukemia (ALL) through analysis of rearrangements in the IGH gene to estimate Minimal Residual Disease (MRD) levels. <input type="checkbox"/> For diagnostic evaluation of rearrangements in the IGH gene in patients with suspected B-cell Non-Hodgkin's Lymphoma (NHL), but in whom clinical, immunophenotypic, and histologic evaluation have provided inconclusive results. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> IGK	<input type="checkbox"/> For medical management of patients with ALL through analysis of rearrangements in the IGK gene to estimate MRD levels. <input type="checkbox"/> For diagnostic evaluation of rearrangements in the IGK gene in patients with suspected B-cell NHL, but in whom clinical, immunophenotypic, and histologic evaluations have provided inconclusive results. <input type="checkbox"/> Other indication		
<input type="checkbox"/> IL28B	<input type="checkbox"/> For IL28B single nucleotide polymorphism (SNP) testing in patients with chronic Hepatitis C Virus (HCV) genotype 1 being considered for treatment with PegIFN/RBV dual therapy. <input type="checkbox"/> Other indication		
<input type="checkbox"/> JAK2	<input type="checkbox"/> Diagnostic evaluation of individuals presenting with clinical, laboratory, or pathological findings suggesting classic forms of myeloproliferative neoplasms (MPN), that is, Polycythemia Vera (PV), Essential Thrombocythemia (ET), or Primary Myelofibrosis (PMF). <input type="checkbox"/> Diagnostic evaluation of PV through JAK2 Exon 12 variant detection in JAK2 p.Val617Phe negative individuals. <input type="checkbox"/> Other indication		
<input type="checkbox"/> KCNQ1, KCNH2, SCN5A, KCNE1, and/or KCNE2	<input type="checkbox"/> For patients with suspected familial Long QT Syndrome for confirmation of diagnosis and treatment. <input type="checkbox"/> Other indication		
<input type="checkbox"/> KIT	<input type="checkbox"/> To confirm a diagnosis of a gastrointestinal stromal tumor (GIST) in patients who are negative by immunostaining. <input type="checkbox"/> To determine primary resistance to treatment with TKIs in patients with an advanced metastatic or unresectable GIST. <input type="checkbox"/> To determine primary resistance to preoperative or postoperative treatment of a GIST with TKIs. <input type="checkbox"/> Other indication		
<input type="checkbox"/> KMT2D and/or KDM6A	<input type="checkbox"/> To confirm a diagnosis of Kabuki Syndrome (KS) in patients with symptoms compatible with KS. <input type="checkbox"/> Other indication		
<input type="checkbox"/> KRAS	<input type="checkbox"/> To help guide administration of anti-EGFR monoclonal antibodies. <input type="checkbox"/> Other indication		
<input type="checkbox"/> MECP2	<input type="checkbox"/> Testing for MECP2 sequence variants in patients who meet established clinical diagnostic criteria for classic or variant Rett Syndrome (RS). <input type="checkbox"/> Testing for MECP2 sequence variants in patients who have symptoms of RS, but do not meet established clinical diagnostic criteria. <input type="checkbox"/> Other indication		
<input type="checkbox"/> MEFV	<input type="checkbox"/> In patients exhibiting symptoms of Familial Mediterranean Fever (FMF), including periodic episodes of fever in combination with peritonitis, pleuritic, arthritis, and erysipelas-like erythema. <input type="checkbox"/> In patients from ethnic groups considered at high risk for FMF who present with nephrotic syndrome or amyloidosis, but do not meet the diagnostic criteria for FMF. <input type="checkbox"/> Other indication		
<input type="checkbox"/> MLH1, MSH2, MSH6, MSI, PMS2, and/or EPCAM	Please provide the clinical indication(s) for genetic testing for Lynch Syndrome (LS)/MLH1, MSH2, MSH6, MSI, PMS2, and/or EPCAM:		

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<input type="checkbox"/> MPL	<input type="checkbox"/> Diagnostic evaluation of Myeloproliferative Leukemia (MPL) variants to include Trp515Leu and Trp515Lys in JAK2 p.Val617Phe-negative individuals showing symptoms. <input type="checkbox"/> Other indication		
<input type="checkbox"/> MUTYH	<input type="checkbox"/> Diagnosis of MYH-Associated Polyposis (MAP) in APC-negative colorectal polyposis patients, or in polyposis patients who have a family history consistent with autosomal recessive inheritance. <input type="checkbox"/> Diagnosis of MAP in asymptomatic siblings of patients with known MYH variants. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Noninvasive Prenatal Screening for Trisomies 13, 18, 21, X & Y	<input type="checkbox"/> In singleton pregnancies with a high risk of fetal aneuploidy. <input type="checkbox"/> Other indication Note, the high-risk criteria is as follows: – Maternal age 35 years or older at delivery – Sonographic findings indicating an increased risk of aneuploidy – History of a prior pregnancy with a trisomy – Positive screening results for aneuploidy, including first trimester, sequential, integrated, or quadruple screen – Parental balanced Robertsonian translocation with increased risk for trisomy 13 or 21		
<input type="checkbox"/> NPM1	<input type="checkbox"/> To guide treatment decisions for individuals with AML. <input type="checkbox"/> Other indication		
<input type="checkbox"/> NRAS	<input type="checkbox"/> For patients with metastatic colorectal cancer who are being considered for treatment with anti-EGFR monoclonal antibodies, and who have had negative KRAS gene testing. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Oncotype DX® Breast Cancer Assay (Oncotype DX®)	<input type="checkbox"/> Estrogen Receptor (ER) positive (+), lymph node (LN) negative (-), human EGFR 2 negative (HER2-) breast cancer patients who are considering whether to use adjuvant chemotherapy in addition to standard hormone therapy. <input type="checkbox"/> ER+, HER2- breast cancer patients with 1-3 involved ipsilateral axillary lymph nodes who are considering whether to use adjuvant chemotherapy in addition to hormonal therapy. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PAX8	<input type="checkbox"/> For individuals with indeterminate thyroid FNA biopsy cytology for diagnosis of papillary thyroid carcinoma. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PDGFRA	<input type="checkbox"/> To confirm a diagnosis of a GIST in patients who are negative by immunostaining. <input type="checkbox"/> To determine primary resistance to treatment with TKIs in patients with an advanced metastatic or unresectable GIST. <input type="checkbox"/> To determine primary resistance to preoperative or postoperative treatment of a GIST with TKIs. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PML/RARalpha	<input type="checkbox"/> Diagnostic assessment of individuals with suspected acute promyelocytic leukemia (APL) by quantitative RT-PCR (RQ-PCR). <input type="checkbox"/> Diagnostic assessment of individuals with suspected APL by qualitative RT-PCR. <input type="checkbox"/> Monitoring response to treatment and disease progression in individuals with APL by RQ-PCR. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> PMP22	<input type="checkbox"/> For the accurate diagnosis and classification of hereditary polyneuropathies. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PPP2R2B	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 12 (SCA12) in patients with action tremor of the upper extremities and signs of cerebellar and cortical dysfunction, in addition to Indian ancestry and a family history consistent with autosomal dominant inheritance. <input type="checkbox"/> Diagnosis of SCA12 in symptomatic family members of known SCA12 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PRSS1	<input type="checkbox"/> A family history of pancreatitis in a first-degree (parent, sibling, child) or second-degree (aunt, uncle, grandparent) relative; <input type="checkbox"/> An unexplained episode of documented pancreatitis occurring in a child that has required hospitalization, and where there is significant concern that hereditary pancreatitis should be excluded; <input type="checkbox"/> Recurrent (two or more separate, documented episodes with hyper-amylasemia) attacks of acute pancreatitis for which there is no explanation (anatomical anomalies, ampullary or main pancreatic strictures, trauma, viral infection, gallstones, alcohol, drugs, hyperlipidemia, etc.); or <input type="checkbox"/> Unexplained (idiopathic) chronic pancreatitis. <input type="checkbox"/> Other indication		
<input type="checkbox"/> PTEN	<input type="checkbox"/> For patients with ASDs and macrocephaly (Head circumference greater than 2 standard above the mean for age). <input type="checkbox"/> PTEN variant testing in individuals suspected of being affected with Cowden Syndrome (CS) or Bannayan-Riley-Ruvalcaba Syndrome (BRRS). <input type="checkbox"/> Other indication		
<input type="checkbox"/> RET	<input type="checkbox"/> Multiple endocrine neoplasia type 2 (MEN2) gene testing in patients with the clinical manifestations of MEN2A, MEN2B, or familial medullary thyroid carcinoma (FMTC), including those with apparently sporadic Medullary Thyroid Carcinoma (MTC) or pheochromocytoma. <input type="checkbox"/> MEN2 gene testing to confirm a diagnosis in the at-risk relatives of genetically confirmed MEN2 patients. <input type="checkbox"/> Other indication		
<input type="checkbox"/> ROS1	<input type="checkbox"/> For patients who have wild type (negative) EGFR or ALK gene testing, reflex testing to ROS1 should be ordered for the treatment of non-small cell lung carcinoma. <input type="checkbox"/> Other indication		
<input type="checkbox"/> RYR1	<input type="checkbox"/> To test clinically confirmed Malignant Hyperthermia Susceptibility (MHS) patients for variants in the RYR1 gene to facilitate diagnostic testing in at-risk relatives. <input type="checkbox"/> To diagnose MHS in at-risk relatives of patients with clinically confirmed MHS. <input type="checkbox"/> Other indication		
<input type="checkbox"/> SDHA, SDHB, SDHC, SDHD, SDHAF2, MAX, and/or TMEM127	<input type="checkbox"/> To diagnose a hereditary paraganglioma (PGL) or pheochromocytoma (PCC) syndrome in patients with PGLs and/or PCCs. <input type="checkbox"/> Other indication		
<input type="checkbox"/> SERPINA1	<input type="checkbox"/> For guidance in diagnosis of inconclusive cases of Alpha-1 Antitrypsin Deficiency (AATD) in individuals with Chronic Obstructive Pulmonary Disease (COPD), unexplained liver disease, family history of AATD, or environmental exposures leading to airflow obstruction after serum Alpha-1 Antitrypsin (AAT) protein levels and protein phenotyping has been completed. <input type="checkbox"/> Other indication		

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<input type="checkbox"/> SMAD4	<input type="checkbox"/> To clarify the diagnosis of individuals with JPS. <input type="checkbox"/> If a known SMAD4 mutation is in the family, genetic testing should be performed in the first six months of life due to hereditary hemorrhagic telangiectasia risk. <input type="checkbox"/> Other indication		
<input type="checkbox"/> SMN1/SMN2	<input type="checkbox"/> Diagnosis of patients with hypotonia and muscle weakness who are suspected of having Spinal Muscular Atrophy (SMA). <input type="checkbox"/> Other indication		
<input type="checkbox"/> SNRPN/UBE3A	When a clinical diagnosis of Prader-Willi Syndrome (PWS) is suspected, the following findings justify genetic testing: <input type="checkbox"/> From birth to age two: Hypotonia with poor suck (neonatal period). <input type="checkbox"/> From age two to age six: Hypotonia with history of poor suck, global developmental delay. <input type="checkbox"/> From age six to age 12: Hypotonia with history of poor suck, global developmental delay, excessive eating with central obesity if uncontrolled. <input type="checkbox"/> From age 13 years to adulthood: Cognitive impairment, usually mild intellectual disability; excessive eating with central obesity if uncontrolled, hypothalamic hypogonadism and/or typical behavior problems. <input type="checkbox"/> Other indication		
	When a clinical diagnosis of Angelman Syndrome is suspected, the following findings justify genetic testing: <input type="checkbox"/> As part of the evaluation of patients with developmental delay, regardless of age. <input type="checkbox"/> As part of the evaluation of patients with a balance or movement disorder such as ataxia of gait. May not appear as frank ataxia but can be forward lurching, unsteadiness, clumsiness, or quick, jerky motions. <input type="checkbox"/> As part of the evaluation of patients with uniqueness of behavior: any combination of frequent laughter/smiling; apparent happy demeanor; easily excitable personality, often with uplifted hand-flapping or waving movements; hypermotoric behavior. <input type="checkbox"/> Speech impairment, none or minimal use of words; receptive and non-verbal communication skills higher than verbal ones. <input type="checkbox"/> Other indication		
<input type="checkbox"/> STK11	<input type="checkbox"/> To confirm a diagnosis of Peutz-Jeghers Syndrome (PJS) in proband patients with a presumptive or probable diagnosis of PJS. <input type="checkbox"/> Other indication		
<input type="checkbox"/> TBP	<input type="checkbox"/> Diagnosis of Spinocerebellar Ataxia Type 17 (SCA17) in ataxia patients exhibiting variable combinations of cognitive decline, psychiatric disturbance, and movement disorders. <input type="checkbox"/> Diagnosis of SCA17 in symptomatic family members of known SCA17 patients. <input type="checkbox"/> Diagnosis of SCA17 in patients suspected of having Huntington Disease (HD) who have tested negative for a pathogenic variant in the HD gene. <input type="checkbox"/> Other indication		
<input type="checkbox"/> TGFBR2	<input type="checkbox"/> To facilitate the diagnosis of Marfan syndrome in patients testing negative for FBN1 gene variants. <input type="checkbox"/> Other indication		
<input type="checkbox"/> TP53	<input type="checkbox"/> Diagnosis of patients satisfying the criteria for classic Li-Fraumeni Syndrome (LFS) or Li-Fraumeni-Like Syndrome (LFLS), or the Chompret criteria for TP53 gene testing. <input type="checkbox"/> Other indication		

COLUMN I Select the gene(s) being requested	COLUMN II Select the indication(s) for the requested test	COLUMN III Indicate the CPT® code(s) and quantity	
		CPT® code(s)	QTY
<input type="checkbox"/> TPMT	<input type="checkbox"/> TPMT genotyping or phenotyping in patients with Inflammatory Bowel Disease (IBD) prior to administration of thiopurines (azathioprine, 6-MP, and 6-TG). <input type="checkbox"/> Other indication		
<input type="checkbox"/> TRG	<input type="checkbox"/> Diagnosis and treatment of T-cell neoplasms. <input type="checkbox"/> Other indication		
<input type="checkbox"/> UGT1A1	<input type="checkbox"/> Prior to irinotecan administration in patients with CRC to lower the starting dose of irinotecan in patients with the UGT1A1*28/UGT1A1*28 genotype. <input type="checkbox"/> Prior to irinotecan administration in patients with CRC to increase the starting dose of irinotecan in patients with the UGT1A1*1/UGT1A1*1 or UGT1A1*1/UGT1A1*28 genotypes. <input type="checkbox"/> Other indication		
<input type="checkbox"/> UPD	<input type="checkbox"/> For neonates, infants, children or adults symptomatic for Beckwith-Wiedemann Syndrome (BWS) to diagnose Uniparental Disomy (UPD) for chromosome 11. <input type="checkbox"/> Other indication		
<input type="checkbox"/> VHL	<input type="checkbox"/> Diagnosis of Von Hippel-Lindau (VHL) syndrome in patients presenting with pheochromocytoma, paraganglioma, or central nervous system hemangioblastoma. <input type="checkbox"/> Confirmation of diagnosis in individuals with symptoms consistent with VHL syndrome. <input type="checkbox"/> Other indication		
<input type="checkbox"/> VKORC1	<input type="checkbox"/> For the initiation and management of warfarin treatment. <input type="checkbox"/> Other indication		
<input type="checkbox"/> Y Chromosome Microdeletion Analysis	<input type="checkbox"/> For detecting submicroscopic deletions involving the Y chromosome in men with azoospermia, oligozoospermia, or teratozoospermia. <input type="checkbox"/> Other indication		

SECTION II – Laboratory developed tests that are NOT covered under the DHA Evaluation of Non-United States FDA Approved LDT Demonstration Project (test/gene not listed in Section I)

Please list the exact genetic test name, CPT® code(s), FDA approval status of the test, and the name of the laboratory performing the test.

Genetic Test Name:

CPT® codes:

Is this an FDA-approved test? Visit www.accessdata.fda.gov/scripts/cdrh/devicesatfda/index.cfm to verify.

Yes No Unknown

Which laboratory is performing the genetic test?

I attest the information provided is true and accurate to the best of my knowledge. I understand Health Net Federal Services, LLC or designee may perform a routine audit and request the medical documentation to verify the accuracy of the information reported on this form.

Additional information: _____

Physician's printed name and title: _____

TIN: _____

Physician signature: _____ Date: _____

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