Introduction

Connective tissue is one kind of tissue that is found in the body. It connects and provides support to other tissues such as muscles, nerves, and the skin. For example, fat, bone and cartilage are types of connective tissues. Some problems with connective tissue can be inherited. This policy describes when it may be medically necessary to do genetic testing to look for inherited connective tissue disorders.

Note: The Introduction section is for your general knowledge and is not to be taken as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals. It is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider also can be a place where medical care is given, like a hospital, clinic, or lab. This policy informs them about when a service may be covered.

Policy Coverage Criteria

<table>
<thead>
<tr>
<th>Testing</th>
<th>Medical Necessity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Individual gene variant testing</td>
<td>Individual genetic testing for the diagnosis of Marfan syndrome, other syndromes associated with thoracic aortic</td>
</tr>
</tbody>
</table>
### Testing

<table>
<thead>
<tr>
<th>Medical Necessity</th>
</tr>
</thead>
<tbody>
<tr>
<td>aneurysms and dissections, and related disorders may be considered medically necessary when:</td>
</tr>
<tr>
<td>- Using established diagnostic criteria the signs and symptoms of a connective tissue disorder are present, but a definitive diagnosis cannot be made without genetic testing.</td>
</tr>
<tr>
<td><strong>AND</strong></td>
</tr>
<tr>
<td>- The focused variant testing panel is limited to the following genes: ACTA2, TGFBR1, TGFBR2 (CPT 81405) and/or FBN1 and MYH11 (CPT 81408).</td>
</tr>
<tr>
<td>Targeted familial variant testing may sometimes be considered medically necessary in asymptomatic individuals in order to assess future risk of disease. This may be the case when there is a known pathogenic variant in the family for Marfan syndrome, other syndromes associated with thoracic aortic aneurysms and dissections, and related disorders. (See Additional Information section below)</td>
</tr>
</tbody>
</table>

### Investigational

<table>
<thead>
<tr>
<th>Panel testing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Genetic testing panels that do not meet the criteria for limited focused gene variant testing described above are considered investigational. (See Additional Information section below)</td>
</tr>
</tbody>
</table>

### Additional Information

- Tissues that surround organs, blood vessels, and bones are called connective tissue. Changes to certain genes may cause problems with connective tissue. Specific genes can be tested to diagnose connective tissue problems.
- Syndromes associated with thoracic aortic aneurysms may have established clinical criteria with major and minor criteria, eg, Marfan syndrome (Ghent criteria) and Ehlers-Danlos syndrome type IV, or may be associated with characteristic clinical findings. While most of these syndromes can be diagnosed based on clinical findings, these syndromes may be associated with variability in clinical presentation and may show overlapping features with each other, and with other disorders. The use of genetic testing to establish a diagnosis in a patient with a suspected connective tissue disorder is most useful in those patients who do not meet sufficient clinical diagnostic criteria at the time of initial examination, in patients who have an
**Additional Information**

*atypical phenotype and other connective tissue disorders cannot be ruled out, and in individuals who belong to a family in which a pathogenic variant is known (presymptomatic diagnosis).*

- Genetic testing has conventionally been used when a definitive diagnosis of one of these syndromes cannot be made. More recently, panels using next-generation sequencing (NGS), which test for multiple genes simultaneously, have been developed for the syndromes associated with thoracic aortic aneurysms and dissections, and other conditions that may have overlapping phenotypes. Although the laboratory-reported sensitivity is high for some of the conditions on the panel, the analytic validity of these panels is unknown, and detection rates of variants of uncertain significance are unknown.

- However, there may be certain clinical scenarios in which focused panel testing may be appropriate to include a narrow list of differential diagnoses of thoracic aortic aneurysms and dissection based on clinical findings.

**Coding**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CPT</td>
<td></td>
</tr>
<tr>
<td>81401</td>
<td>MED12 (mediator complex subunit 12)(eg, FG syndrome type 1, Lujan syndrome), common variants (eg, R961W, N1007S)</td>
</tr>
<tr>
<td>81405</td>
<td>ACTA2 (actin, alpha 2, smooth muscle, aorta) (eg, thoracic aortic aneurysms and aortic dissections), full gene sequence TGFBR1 (transforming growth factor, beta receptor 1) (eg, Marfan syndrome), full gene sequence</td>
</tr>
<tr>
<td>81408</td>
<td>FBN1 (fibrillin 1) (eg, Marfan syndrome), full gene sequence MYH11 (myosin, heavy chain 11, smooth muscle) (eg, thoracic aortic aneurysms and aortic dissections), full gene sequence</td>
</tr>
<tr>
<td>81410</td>
<td>Aortic dysfunction or dilation (eg, Marfan syndrome, Loeys Dietz syndrome, Ehlers-Danlos syndrome type IV, arterial tortuosity syndrome); genomic sequence analysis panel, must include sequencing of at least 9 genes, including FBN1, TGFBR1, TGFBR2, COL3A1, MYH11, ACTA2, SLC2A10, SMAD3, and MYLK</td>
</tr>
<tr>
<td>81411</td>
<td>Aortic dysfunction or dilation (eg, Marfan syndrome, Loeys Dietz syndrome, Ehler Danlos syndrome type IV, arterial tortuosity syndrome); duplication/deletion analysis panel, must include analyses for TGFBR1, TGFBR2, MYH11, and COL3A1</td>
</tr>
<tr>
<td>Code</td>
<td>Description</td>
</tr>
<tr>
<td>-------</td>
<td>---------------------------------</td>
</tr>
<tr>
<td>81479</td>
<td>Unlisted molecular pathology code</td>
</tr>
</tbody>
</table>

**Note:** CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). HCPCS codes, descriptions and materials are copyrighted by Centers for Medicare Services (CMS).

## Individual Gene Testing

For individual gene testing, the CPT codes in **Table 1** may be used.

### Table 1: Coding for Individual Gene Testing

<table>
<thead>
<tr>
<th>Disease</th>
<th>Associated Gene</th>
<th>Percentage of Probands With a Pathogenic Variant Detected by Method</th>
<th>CPT Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Diseases associated with TAAD</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Marfan syndrome</td>
<td>FBN1</td>
<td>Sequence analysis detects 70%-93%</td>
<td>Included in 81408</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Deletion/duplication analysis detection rate unknown</td>
<td></td>
</tr>
<tr>
<td>EDS type IV (vascular type)</td>
<td>COL3A1</td>
<td>Sequence analysis detects &gt;95%</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Deletion/duplication analysis detects ≈2%</td>
<td></td>
</tr>
<tr>
<td>LDS</td>
<td>TGFBR1</td>
<td>Percentage of LDS attributed to variants in following genes by sequence analysis:</td>
<td>TGFBR1: in 81405</td>
</tr>
<tr>
<td></td>
<td>TGFBR2</td>
<td>TGFBR1: 20%</td>
<td>TGFBR2: in 81405</td>
</tr>
<tr>
<td></td>
<td>SMAD3</td>
<td>TGFBR2: 70%</td>
<td>SMAD3 and TGFBR2: unlisted 81479</td>
</tr>
<tr>
<td></td>
<td>TGFB2</td>
<td>SMAD3: 5%</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>TGFB2: 1%</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>In general, variants detected in LDS by deletion/duplication analysis are not associated with aortic aneurysms</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Familial TAAD</td>
<td>TGFBR1</td>
<td>Percentage of familial TAAD attributed to variants in following genes by sequence and deletion/duplication analysis:</td>
<td>TGFBR1: in 81405</td>
</tr>
<tr>
<td></td>
<td>TGFBR2</td>
<td>TGFBR1: 1%</td>
<td>TGFBR2: in 81405</td>
</tr>
<tr>
<td></td>
<td>MYH11</td>
<td>TGFBR2: 4%</td>
<td>MYH11: in 81408</td>
</tr>
<tr>
<td></td>
<td>ACTA2</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>FBN1</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>MYLK</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>Disease</td>
<td>Associated Gene</td>
<td>Percentage of Probands With a Pathogenic Variant Detected by Method</td>
<td>CPT Codes</td>
</tr>
<tr>
<td>-----------------------------------------------------------</td>
<td>-----------------</td>
<td>-------------------------------------------------------------------------------------------------------------------------------------</td>
<td>--------------------</td>
</tr>
<tr>
<td>SMAD3</td>
<td></td>
<td>ACTA2: 10%-14%</td>
<td>unlisted 81479</td>
</tr>
<tr>
<td></td>
<td></td>
<td>FBN1: unknown</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Sequence analysis:</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>MYLK: 1%</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>SMAD3: 2%</td>
<td></td>
</tr>
<tr>
<td>Arterial tortuosity syndrome</td>
<td>SLC2A10</td>
<td>Sequence analysis detects ≈86%</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Deletion/duplication analysis detects ≈7%</td>
<td></td>
</tr>
</tbody>
</table>

**Diseases not associated with TAAD**

<table>
<thead>
<tr>
<th>MED12-related disorders (FGS syndrome type 1 and Lujan syndrome)</th>
<th>MED12</th>
<th>Variant detection frequency unknown</th>
<th>Included in 81401</th>
</tr>
</thead>
<tbody>
<tr>
<td>Shprintzen-Goldberg syndrome</td>
<td>SK1</td>
<td>Sequence analysis and deletion/duplication analysis rates of detection have not been reported</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td>EDS classic type (EDS I and II)</td>
<td>COL5A1 COL5A2</td>
<td>Percentage of EDS classic type attributed to variants in following genes by sequence analysis: COL5A1: 46% COL5A2: 4%</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td>EDS kyphoscoliotic form (EDS type VI)</td>
<td>PLOD1</td>
<td>Variant detection frequency by sequence analysis is unknown  Deletions/duplications have been detected with a frequency of 18%</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td>Periventricular heterotopia, EDS variant</td>
<td>FLNA</td>
<td>Sequence analysis 100% in those with family history and 26% in simplex females  Detection by deletion/duplication analysis is unknown</td>
<td>Unlisted 81479</td>
</tr>
<tr>
<td>Congenital contractual arachnodactyly</td>
<td>FBN2</td>
<td>Sequence analysis has been reported to detect 27%-75% of variants  Variant detection by deletion/duplication analysis is unknown</td>
<td>Unlisted 81479</td>
</tr>
</tbody>
</table>

- EDS classic type.
- Loeys-Dietz syndrome.
- TAAD
Panel Testing

Specific CPT codes for genetic panel tests associated with aortic dysfunction or dilation syndromes (81410 and 81411) are described in the coding table above with the genes included in each test.

Related Information

Genetic Counseling

Genetic counseling is primarily aimed at patients who are at risk for inherited disorders, and experts recommend formal genetic counseling in most cases when genetic testing for an inherited condition is considered. The interpretation of the results of genetic tests and the understanding of risk factors can be very difficult and complex. Therefore, genetic counseling will assist individuals in understanding the possible benefits and harms of genetic testing, including the possible impact of the information on the individual's family. Genetic counseling may alter the utilization of genetic testing substantially and may reduce inappropriate testing. Genetic counseling should be performed by an individual with experience and expertise in genetic medicine and genetic testing methods.

Evidence Review

Description

Marfan syndrome (MFS) is a systemic connective tissue disorder that may have a high degree of clinical variability and overlapping phenotypes with other syndromes and disorders. The diagnosis of most suspected connective tissue disorders can be made based on clinical findings and family history. Some of these disorders are associated with a predisposition to the development of progressive thoracic aortic aneurysms and dissection (TAAD). Accurate diagnosis of one of these syndromes can lead to changes in clinical management, including surveillance of the aorta and surgical repair of the aorta, when necessary, as well as surveillance for multisystem involvement in syndromic forms of TAAD. Known genetic variants are associated with MFS and the other connective tissue disorders that may share clinical features with MFS.
Background

Connective Tissue Diseases

Individuals suspected of having a systemic connective tissue disorder like MFS usually have multiple features that affect many different organ systems, and most of these conditions can be diagnosed using clinical criteria. However, these different syndromes may show shared features, overlapping phenotypes, and similar inheritance patterns, which can cause a diagnostic challenge. Additional difficulties in the diagnosis of one of these syndromes may occur due to the age-dependent development of many of the physical manifestations of the syndrome (making the diagnosis more difficult in children). Many syndromes show variable expression, and numerous features found in many of these syndromes occur in the general population (eg, pectus excavatum, tall stature, joint hypermobility, mitral valve prolapse, nearsightedness). The identification of the proper syndrome is important to address its manifestations and complications, in particular, the risk of aortic aneurysms and dissection.

Thoracic Aortic Aneurysms and Dissection

Most TAAs are degenerative and are often associated with the same risk factors as abdominal aortic aneurysms (eg, atherosclerosis). TAAs may be associated with a genetic predisposition, which can either be familial or related to defined genetic disorders or syndromes.1

Genetic predisposition to TAA is due to a genetic defect that leads to abnormalities in connective tissue metabolism. Genetically-related TAA accounts for approximately 5% of TAA.1 Some of the genetic syndromes associated with TAA have more aggressive rates of aortic expansion and are more likely to require intervention compared with sporadic TAA. MFS is the most common inherited form of syndromic TAA and TAAD. Other genetic systemic connective tissue disorders associated with a risk of TAAD include Ehlers-Danlos syndrome (EDS) type IV, Loeys-Dietz syndrome (LDS), and arterial tortuosity syndrome.

Familial TAAD refers to patients with a family history of aneurysmal disease, but who do not meet criteria for a connective tissue syndrome.
Marfan Syndrome

Marfan Syndrome (MFS) is an autosomal dominant condition. There is a high degree of clinical variability of systemic manifestations, ranging from isolated features of MFS to neonatal presentation of severe and rapidly progressive disease in multiple organ systems. Despite the clinical variability, the principal manifestations involve the skeletal, ocular and cardiovascular systems. Involvement of the skeletal system is characterized by bone overgrowth and joint laxity, disproportionately long extremities for the size of the trunk (dolichostenomelia), overgrowth of the ribs which can push the sternum in or out (pectus excavatum or carinatum, respectively), and scoliosis which can be mild or severe and progressive. Ocular features include myopia, and displacement of the lens from the center of the pupil (ectopic lentis) is a hallmark feature and is seen in 60% of affected individuals. Cardiovascular manifestations are the major source of morbidity and mortality and include dilation of the aorta at the level of the sinuses of Valsalva, predisposition for aortic tear and rupture, mitral valve prolapse, tricuspid valve prolapse and enlargement of the proximal pulmonary artery. However, with proper management, the life expectancy of someone with MFS can approximate that of the general population.

The diagnosis of MFS is mainly a clinical one and based on the characteristic findings in multiple organ systems, as well as the family history. The Ghent criteria, revised in 2010, are used for the clinical diagnosis of MFS. The previous Ghent criteria had been criticized for taking insufficient account of the age-dependent nature of some of the clinical manifestations, making the diagnosis in children more difficult. They were also criticized for including some non-specific physical manifestations or poorly validated diagnostic thresholds. The revised criteria were based on clinical characteristics in large published patient cohorts, as well as expert opinions of panel members with extensive experience in application of the criteria, the differential diagnosis of MFS, and the strengths and limitations of molecular genetic testing. The revised criteria have 5 major changes to the previous diagnostic guidelines. First, more weight is given to the 2 cardinal features of MFS, aortic root aneurysm/dissection and ectopic lentis. In the absence of findings that are not expected in MFS, the combination of these 2 features is sufficient to make the diagnosis. When aortic disease is present, but ectopia lentis is not, all other cardiovascular and ocular manifestations of MFS and findings in other organ systems contribute to a “systemic score” that guides diagnosis. Second, a more prominent role has been given to molecular testing of FBN1 and other relevant genes, allowing for the appropriate use when necessary. Third, some of the less specific manifestations of MFS were removed or made less influential in the diagnostic criteria. Fourth, the revised criteria formalize the concept that additional diagnostic considerations and testing may be required if a patient has findings that satisfy the criteria for MFS but show unexpected findings, particularly if they are suggestive of a specific alternative diagnosis. Particular emphasis is placed on LDS, Sphrintzen-Goldberg syndrome...
(SGS), and EDS-vascular type. LDS and SGS may have substantial overlap with MFS, including the potential for similar involvement of the aortic root, skeleton, skin and dura. EDS-vascular type occasionally shows overlap with MFS. Each of these conditions has a unique risk profile and management protocol.\textsuperscript{3} Given the autosomal dominant inheritance, the number of physical findings needed to establish a diagnosis for someone with an established family history is reduced.

It is estimated that molecular techniques allow for the detection of FBN1 mutations in up to 97% of Marfan patients who fulfil Ghent criteria, suggesting that the current Ghent criteria have excellent specificity.\textsuperscript{3}

FBN1 is the only gene in which mutations are known to cause classic MFS. Approximately 75% of individuals with MFS have an affected parent, and 25% have a de novo mutation. Over 1000 FBN1 mutations that cause MFS have been identified. The following findings in FBN1 molecular genetic testing should infer causality in making the diagnosis of MFS: a pathogenic variant previously shown to segregate in families with MFS and de novo mutations of a certain type (eg, nonsense, certain missense mutations, certain splice site mutations, certain deletions and insertions).\textsuperscript{2}

Most mutations in the FBN1 gene that cause MFS can be identified with sequence analysis (≈70\%-93\%) and, although the yield of deletion/duplication analysis in patients without a defined coding sequence or splice site by sequence analysis is unknown, it is estimated to be about 30\%. The most common testing strategy of a proband suspected of having MFS is sequence analysis followed by deletion/duplication analysis if a pathogenic variant is not identified.\textsuperscript{2} However, the use of genetic testing for a diagnosis of MFS has limitations. More than 90\% of mutations that have been described are unique, and most mutations are not repeated among non-genetically related patients. Therefore, the absence of a known mutation in a patient in whom MFS is suspected does not exclude the possibility that the patient has MFS. No clear genotype-phenotype correlation exists for MFS and, therefore the severity of the disease cannot be predicted from the type of mutation.

Caution should be used in interpreting the identification of a FBN1 mutation, as other conditions with overlapping phenotypes with MFS can have an FBN1 mutation (eg, MASS syndrome, familial mitral valve prolapse syndrome, SGS, isolated ectopic lentis).

Management of MFS includes both treatment of manifestations and prevention of complications, including surgical repair of the aorta depending on the maximal measurement, the rate of increase of the aortic root diameter, and the presence of progressive and severe aortic regurgitation.
**Ehlers-Danlos Syndrome**

Ehlers-Danlos Syndrome (EDS) is a group of disorders that affect connective tissues and share common features characterized by skin hyperelasticity or laxity, abnormal wound healing, and joint hypermobility. The defects in connective tissues can vary from mildly loose joints to life-threatening complications. All types of EDS affect the joints and many affect the skin, but features vary by type.

The different types of EDS include types I and II (classic type), type III (hypermobility type), type IV (vascular type), type VI (kyphoscoliotic form), all of which are inherited in an autosomal dominant pattern with the exception of type VI, which is autosomal recessive. It is estimated that affected individuals with types I, II or IV may inherit the disease-causing variant from an affected parent 50% of the time, and about 50% have a de novo disease-causing variant.

Most types of EDS are not associated with aortic dilation, with the exception of the vascular type (also known as type IV), which can involve serious and potentially life-threatening complications. The prevalence of the vascular type may affect about 1 in 250,000 people. Vascular complications include rupture, aneurysm, and/or dissection of major or minor arteries. Arterial rupture may be preceded by aneurysm, arteriovenous fistulae or dissection, or may occur spontaneously. Such complications are often unexpected and may present as sudden death, stroke, internal bleeding and/or shock. The vascular type is also associated with an increased risk of gastrointestinal perforation or organ rupture, and rupture of the uterus during pregnancy.

The clinical diagnosis of EDS type IV can be made from major and minor clinical criteria. The combination of 2 major criteria (arterial rupture, intestinal rupture, uterine rupture during pregnancy and a family history of EDS type IV) is highly specific. The presence of one or more minor clinical criteria supports the diagnosis, but is insufficient to make the diagnosis by itself.

Pathogenic variants in the COL1A1, COL1A2, COL3A1, COL5A1, COL5A2, PLOD1, and TNXB genes cause EDS. The vascular type (type IV) is caused by pathogenic variants in the COL3A1 gene.

**Loeys-Dietz Syndrome**

Loeys-Dietz Syndrome (LDS) is an autosomal dominant condition that is characterized by 4 major groups of clinical findings, including vascular, skeletal, craniofacial, and cutaneous manifestations. Vascular findings include cerebral, thoracic, and abdominal arterial aneurysms.
and/or dissections. Skeletal findings include pectus excavatum or carinatum, scoliosis, joint laxity, arachnodactyly and talipes equinovarus. The natural history of LDS is characterized by arterial aneurysms, with a mean age of death of 26 years and a high incidence of pregnancy-related complications, including uterine rupture and death. Treatment considerations take into account that aortic dissection tends to occur at smaller aortic diameters than MFS, and the aorta and its major branches can dissect in the absence of much, if any, dilation. Patients with LDS require echocardiography at frequent intervals to monitor the status of the ascending aorta, and angiography to image the entire arterial tree.

LDS is caused by pathogenic variants in TGFBR1, TGFBR2, TGFB2, and SMAD3 genes.

**Arterial Tortuosity Syndrome**

Arterial tortuosity syndrome is inherited in an autosomal recessive pattern and is characterized by tortuosity of the aorta and/or large- and middle-sized arteries throughout the body. Aortic root dilation, stenosis and aneurysms of large arteries are common. Other features of the syndrome include joint laxity and hyperextensible skin. The syndrome is caused by pathogenic variants in the SLC2A10 gene.

**Familial TAAD**

Approximately 80% of familial TAA and TAAD is inherited in an autosomal-dominant manner and may be associated with variable expression and decreased penetrance of the disease-associated variant.

The major cardiovascular manifestations of familial TAAD (fTAAD) include dilatation of the ascending thoracic aorta at the level of the sinuses of Valsalva or ascending aorta, or both, and dissections of the thoracic aorta involving either the ascending or descending aorta. In the absence of surgical repair of the ascending aorta, affected individuals have progressive enlargement of the ascending aorta, leading to acute aortic dissection. Presentation of the aortic disease and the age of onset are highly variable. fTAAD is diagnosed based on the presence of thoracic aorta pathology, absence of clinical features of MFS, LDS, or vascular EDS, and a positive family history of TAAD. fTAAD is associated with variants in TGFBR2, TGFBR1, MYH11, ACTA2, MYLK, SMAD3, and 2 loci on other chromosomes, AAT1 and AAT2. Rarely, fTAAD can also be caused by FBN1 variants. To date, only about 20% of fTAAD is accounted for by variants in known genes. Early prophylactic repair should be considered in individuals with confirmed
variants in TGFBR2 and TGFBR1 and/or a family history of aortic dissection with minimal aortic enlargement.

The following syndromes and conditions may share some of the features of these connective tissue syndromes, but do not share the risk of TAAD.

**Congenital Contractural Arachnodactyly (Beal Syndrome)**

Congenital contractural arachnodactyly (CCA) is an autosomal dominant condition characterized by a Marfan-like appearance and long, slender toes and fingers. Other features may include “crumpled” ears, contractures of the knees and ankles at birth with improvement over time, camptodactyly, hip contractures, and progressive kyphoscoliosis. Mild dilatation of the aorta is rarely present. CCA is caused by the pathogenic variants in the FBN2 gene.

**MED12-Related Disorders**

The phenotypic spectrum of MED12-related disorders is still being defined, but includes Lujan syndrome (LS) and FG syndrome type 1 (FGS1). LS and FGS1 share the clinical findings of hypotonia, cognitive impairment and abnormalities of the corpus callosum. Individuals with LS share some physical features with MFS, in that they have Marfanoid features including tall and thin habitus, long hands and fingers, pectus excavatum, narrow palate and joint hypermobility. MED12-related disorders are inherited in an X-linked manner, with males being affected and carrier females not usually being affected.

**Shprintzen-Goldberg Syndrome**

Shprintzen-Goldberg Syndrome (SGS) is an autosomal dominant condition that is characterized by a combination of major characteristics which include craniosynostosis, craniofacial findings, skeletal findings, cardiovascular findings, neurologic and brain anomalies, certain radiographic findings and other findings. SK1 is the only gene in which pathogenic variants are known to cause SGS.
**Homocystinuria Caused by Cystathionine Beta-Synthase Deficiency**

Homocystinuria is a rare metabolic disorder inherited in an autosomal recessive manner, which is characterized by an increased concentration of homocysteine, a sulfur-containing amino acid, in the blood and urine. The classical type is due to a deficiency of cystathionine beta synthase (CBS). Affected individuals appear normal at birth but develop serious complications in early childhood, usually by age 3 to 4 years. Heterozygous carriers (1/70 of the general population) have hyperhomocysteinemia without homocystinuria; however, their risk for premature cardiovascular disease is increased.

Overlap with MFS can be extensive and includes a Marfanoid habitus with normal to tall stature, pectus deformity, scoliosis, and ectopia lentis. Central nervous system manifestations include mental retardation, seizures, cerebrovascular events, and psychiatric disorders. Patients have a tendency for intravascular thrombosis and thromboembolic events, which can be life-threatening. Early diagnosis and prophylactic medical and dietary care can decrease and even reverse some of the complications. The diagnosis depends on measurement of CBS activity in tissue (e.g., liver biopsy, skin biopsy).

**Summary of Evidence**

For individuals who have signs and/or symptoms of a connective tissue disease (CTD) linked to thoracic aortic aneurysms who received testing for genes associated with CTDs, the evidence includes mainly clinical validity data. Relevant outcomes are overall survival, disease-specific survival, test accuracy and validity, symptoms, and morbid events. Published data on analytic validity of individual and panel testing of genes is lacking. Sequencing analysis for Marfan syndrome (MFS) has been reported to detect 70% to 93% of pathogenic variants in probands with MFS, and over 95% in Ehlers-Danlos syndrome type IV. Direct evidence of clinical utility is lacking; however, confirming a diagnosis leads to changes in clinical management, which improve health outcomes. These changes in management include treatment of manifestations of a specific syndrome, prevention of primary manifestations and secondary complications, impact on surveillance, and counselling on agents and circumstances to avoid. The evidence is sufficient to determine that the technology results in a meaningful improvement in the net health outcome.

For asymptomatic individuals with a known familial pathogenic variant associated with thoracic aortic aneurysms and dissection who receive targeted familial variant testing, the evidence is generally lacking. Relevant outcomes are overall survival, disease-specific survival, test accuracy and validity, symptoms, and morbid events. Published data on analytic validity of targeted
familial variant testing is lacking, but is expected to be high. Direct evidence of clinical utility is lacking. However, confirming a diagnosis leads to changes in clinical management, which improve health outcomes, similar to those in the proband. In addition, test results will determine whether to follow a relative who does or does not have the familial variant. The evidence is sufficient to determine that the technology results in a meaningful improvement in the net health outcome.

Ongoing and Unpublished Clinical Trials

A search of ClinicalTrials.gov in December 2016 did not identify any ongoing or unpublished trials that would likely influence this policy.

Practice Guidelines and Position Statements

The American College of Medical Genetics

The American College of Medical Genetics issued guidelines on the evaluation of the adolescent or adult with some features of MFS.12

If there is no family history of MFS, then the subject has the condition under any of the following four situations:

1. A dilated aortic root (defined as greater than or equal to two standard deviations above the mean for age, sex, and body surface area) and ectopia lentis

2. A dilated aortic root and a mutation in FBN1 that is clearly pathologic

3. A dilated aortic root and multiple systemic features or

4. Ectopia lentis and a mutation in FBN1 that has previously been associated with aortic disease

If there is a positive family history of MFS (independently ascertained with these criteria), then the subject has the condition under any of the following three situations:

1. Ectopia lentis

2. Multiple systemic features or

3. A dilated aortic root (if over 20 years, greater than two standard deviations; if younger than 20, greater than three standard deviations)
The systemic features are weighted by a scoring system for systemic features.

**Medicare National Coverage**

There is no national coverage determination (NCD). In the absence of an NCD, coverage decisions are left to the discretion of local Medicare carriers.

**Regulatory Status**

Clinical laboratories may develop and validate tests in-house and market them as a laboratory service; laboratory-developed tests (LDTs) must meet the general regulatory standards of the Clinical Laboratory Improvement Amendments (CLIA). Laboratories that offer LDTs must be licensed by CLIA for high-complexity testing. To date, the U.S. Food and Drug Administration has chosen not to require any regulatory review of this test.

Several commercial laboratories currently offer targeted genetic testing, as well as NGS panels that simultaneously analyze multiple genes associated with MFS, TAADs, and related disorders. NGS technology cannot detect large deletions or insertions, and therefore samples that are mutation-negative after sequencing should be evaluated by other testing methodologies.

**Ambry Genetics** offers “TAADNEXT,” an NGS panel that simultaneously analyzes 20 genes that are associated with TAADs, MFS, and related disorders. The panel detects variants in all coding domains and splice junctions of ACTA2, CBS, COL3A1, COL5A1, COL5A2, FBN1, FBN2, FLNA, MED12, MYH11, MYLK, NOTCH1, PLOD1, PRKG1, SKI, SLC2A10, SMAD3, TGFBR2, TGFBR1, and TGFBR2. Gross deletion/duplication analysis is performed for all genes on the panel except CBS, COL5A1, and FLNA.

**GeneDx** offers panel testing “Marfan/TAAD sequencing panel” and “Marfan/TAAD deletion/duplication panel,” which include variant testing for ACTA2, CBS, COL3A1, COL5A1, COL5A2, FBN1, FBN2, FLNA, MED12, MYH11, SKI, SLC2A10, SMAD3, TGFBR2, TGFBR1, and TGFBR2.

**Prevention Genetics** offers targeted familial variant testing, as well as panel testing “Marfan syndrome and related aortopathies next generation sequencing [NGS] panel,” which includes 14 genes: ACTA2, COL3A1, COL5A1, COL5A2, FBN1, FBN2, MYH11, MYLK, SKI, SLC2A10, SMAD3, TGFBR2, TGFBR1, and TGFBR2.
# References


# History

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>05/12/15</td>
<td>New Policy. Policy created with a literature review through November 6, 2014. The use of panels for the detection of mutations in syndromes that may be associated with thoracic aortic aneurysms and dissection is investigational. In certain circumstances, individual mutation testing may be considered medically necessary.</td>
</tr>
<tr>
<td>05/01/16</td>
<td>Annual Review, approved April 12, 2016. Policy updated with a literature review through December 9, 2015; references 10-11 added. Policy statements unchanged.</td>
</tr>
<tr>
<td>09/01/16</td>
<td>Interim Update, approved August 9, 2016. Removed codes 81405 and 81408 from panel testing policy statement.</td>
</tr>
</tbody>
</table>

Policy moved to new format. No changes to policy statement.

**Disclaimer:** This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. The Company adopts policies after careful review of published peer-reviewed scientific literature, national guidelines and local standards of practice. Since medical technology is constantly changing, the Company reserves the right to review and update policies as appropriate. Member contracts differ in their benefits. Always consult the member benefit booklet or contact a member service representative to determine coverage for a specific medical service or supply. CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). ©2017 Premera All Rights Reserved.

**Scope:** Medical policies are systematically developed guidelines that serve as a resource for Company staff when determining coverage for specific medical procedures, drugs or devices. Coverage for medical services is subject to the limits and conditions of the member benefit plan. Members and their providers should consult the member benefit booklet or contact a customer service representative to determine whether there are any benefit limitations applicable to this service or supply. This medical policy does not apply to Medicare Advantage.
Discrimination is Against the Law

Premera Blue Cross complies with applicable Federal civil rights laws and does not discriminate on the basis of race, color, national origin, age, disability, or sex. Premera does not exclude people or treat them differently because of race, color, national origin, age, disability or sex.

Premera:
• Provides free aids and services to people with disabilities to communicate effectively with us, such as:
  • Qualified sign language interpreters
  • Written information in other formats (large print, audio, accessible electronic formats, other formats)
• Provides free language services to people whose primary language is not English, such as:
  • Qualified interpreters
  • Information written in other languages

If you need these services, contact the Civil Rights Coordinator.

If you believe that Premera has failed to provide these services or discriminated in another way on the basis of race, color, national origin, age, disability, or sex, you can file a grievance with:

Civil Rights Coordinator - Complaints and Appeals
PO Box 91102, Seattle, WA 98111
Toll free 855-332-4535, Fax 425-918-5592. TTY 800-842-5357
Email AppealsDepartmentInquiries@Premera.com


Getting Help in Other Languages

This Notice has Important Information. This notice may have important information about your application or coverage through Premera Blue Cross. There may be key dates in this notice. You may need to take action by certain deadlines to keep your health coverage or help with costs. You have the right to get this information and help in your language at no cost. Call 800-722-1471 (TTY: 800-842-5357).

 العربية (Arabic):

يحيى هذه الإشارة معلومات هامة. قد يحيى هذا الإشارة معلومات مهمة مخصصة لطبيك أو
العملية التي لديك الحق في الحصول عليها من خلال Premera Blue Cross. إن تكون هناك إرشادات
في هذا الإشعار. قد تحتاج لإذاعة إجراء في تفويض المدفوع على تكاليف الصحة والرعاية
وفي هذا الكشف. يحمي تلك المعلومات على هذه المعلومات والمساعدة بذلك دون تكيد أي كلمة. يصل
800-722-1471 (TTY: 800-842-5357).

中文 (Chinese):

本通知有重要的讯息。本通知可能有關於您透過 Premera Blue Cross 提交的
申請或保險的重要訊息。本通知可能有重要日期。您可能需要在截止日期
之前採取行動，以保留您的健康保險或費用補貼。您有權利免費以您的母
語得到本訊息和幫助。請接電話 800-722-1471 (TTY: 800-842-5357).

Oromo (Cushite):

Beeksinsi kun odefannoo barbaachisaa qaba. Beeksiisti kun sagantaa
yoorkan karaa Premera Blue Cross tiin tajaajila keessan ilaachiise
odeffan odefannoo barbaachisa qabaachuu danda’a. Guyyaaaw an maraatsaa
ta’an beeksiisaa kan ka seessatti ilaaliia. Tarii kaffaltidhaa deeggarammoo
yoorkan tajaajila fiyyaayaa keessanmif guyyaa dhumaay irratti wanti raawwatan
jaarachuu danda’a. Kaffaltii irraa bilisaa haaltaa ta’an afaan keessanmif
odeffan odefannoo aragchuuf fi deeggarasa aragchuuf miga ni qabaattuu.
Lakkoofsaa biibilaay 800-722-1471 (TTY: 800-842-5357) ti biibilaay.

Français (French):

Cet avis a d’importantes informations. Cet avis peut avoir d’importantes
informations sur votre demande ou la couverture par l’intermédiaire de
Premera Blue Cross. Le présent avis peut contenir des dates clés. Vous
devrez peut-être prendre des mesures par certains délais pour maintenir
votre couverture de santé ou d’aide avec les coûts. Vous avez le droit
d’obtenir cette information et de l’aide dans votre langue à aucun coût.
Appelez le 800-722-1471 (TTY: 800-842-5357).

Kreyól ayisyen (Creole):

Avi si a gen Enfòmasyon Enpòtan Ipadann. Avi si a kapab genyen
enfòmasyon enpòtan konsénan aplikasyon w lan oswa konvènti kouvèti
asinans lan atravè Premera Blue Cross. Kapab genyen dat ki enpòtan nan
avi si a. Ou ka gen pou pran kék aksyon avan sèt man dwa w pou konse kouvèti
asinans sante w la oswa pou yo ka ede w avek depans yo. Se dwa w pou resewwa enfòmasyon sa a ak assisyans nan lang ou pale a, san ou pa gen pou peye pou sa. Rate nan 800-722-1471
(TTY: 800-842-5357).

Deutsche (German):

Diese Benachrichtigung enthält wichtige Informationen. Diese
Benachrichtigung enthält unter Umständen wichtige Informationen
bezüglich Ihres Antrags auf Krankenversicherungsschutz durch Premera
Blue Cross. Suchen Sie nach eventuellen wichtigen Terminen in dieser
Benachrichtigung. Sie könnten bis zu bestimmten Stichtagen handeln
müssen, um Ihren Krankenversicherungsschutz oder Hilfe mit den Kosten
zu behalten. Sie haben das Recht, kostenlos Hilfe und Informationen
in Ihrer Sprache zu erhalten. Rufen Sie an unter 800-722-1471
(TTY: 800-842-5357).

Hmoob (Nmông):

Tsab ntaww tshaj xo no muaj cov ntsiab lus tseem ceeb. Tej zaurn
tsab ntaww tshaj xo no muaj cov ntsiab lus tseem ceeb tsoog kog daim
ntaww thov kev pab los yog kog chov kev pab cuam los ntaww Premera Blue
Cross. Tej zaurn cov hnb tseem ceeb uss sau rau huav daim ntaww no.
Tej zaurn cov kog juaw tau uu gee yam uss peb kom kog uss tis puub
dhuu cov cajj nyong uas teev tseeg rau huav daim ntaww no mas kog kajj
juaw taa baiss kev pab cuam koh moob los yog kev pab them tej nqi kho moob
ntaww. Kog muaj cai kom laww muab cov ntsiab lus no uas tau muab sau
us kog hom lus pub daww rau kog. Hu rau 800-722-1471
(TTY: 800-842-5357).

Ilokko (Ilocano):

Daytoy a Pakkada ket naglao iti Napateg nga Impormasion. Daytoy a
pakkadak balabin nga adda ket naglao iti napateg nga impormasion
maipanggep iti aplikasyonu wno coverage babaen iti Premera Blue
Cross. Daytoy ket malaban dagiti importante a petsa iti daytoy a pakkada.
Balabin nga adda rumbeng nga aramideng nga addang sakkay dagiti
partikular a naituding nga aldaw tapno mapagtalinaedyo ti coverage ti
salay-ano wno tulong kadagit gastos. Adda karbenganyo a mangala iti
daytoy nga impormasion ken tulong ti bukodyo a pagasao nga awan ti

Italiano (Italian):

Questo avviso contiene informazioni importanti. Questo avviso può contenere
informazioni importanti sulla tua domanda o copertura attraverso Premera
Blue Cross. Potrebbero esserci date chiave in questo avviso. Potrebbe
essere necessario un tuo intervento entro una scadenza determinata per
consentirti di mantenere la tua copertura o sovvenzione. Hai il diritto di
ottenere queste informazioni e assistenza nella tua lingua gratuitamente.
Chiama 800-722-1471 (TTY: 800-842-5357).
Este aviso podrá conter informações importantes. Es posible que este aviso contenga información importante acerca de su solicitud o cobertura a través de Premera Blue Cross. Es posible que haya fechas clave en este aviso. Es posible que deba tomar alguna medida antes de determinadas fechas para mantener su cobertura médica o ayuda con los costos. Usted tiene derecho a recibir esta información y ayuda en su idioma sin costo alguno. Llame al 800-722-1471 (TTY: 800-842-5357).

Tagalog (Tagalog):
Ang Paunawa na ito ay naglalaman ng mahalagang impormasyon. Ang paunawa na ito ay naglalaman ng mahalagang impormasyon tungkol sa iyong aplikasyon o pagsakop sa pamagatang ng Premera Blue Cross. Maaaring ma ganang o malalaman mo ang mahalagang impormasyon para sa iyong pagsakop sa kalusugan o tulungan sa iyong kalusugan. Pwede kang iulat o iisipin ang iyong impormasyon sa TTY na 800-722-1471 ng panulad na 800-842-5357.

ไทย (Thai):
ประกาศนี้มีข้อมูลที่สำคัญเกี่ยวกับการเปลี่ยนแปลงหรือการสิ้นสุดสัญญาของคุณใน Premera Blue Cross และมีการแจ้งเตือนในกรณีที่คุณจะต้องให้ความรับผิดชอบในสัญญาของคุณเกี่ยวกับสิทธิของคุณที่สำคัญที่มีผลต่อคุณ ควรติดต่อคุณให้ทราบและแจ้งข้อมูลที่เกี่ยวข้องกับการสิ้นสุดสัญญาของคุณ โปรดติดต่อผู้ให้บริการที่เหมาะสมที่ 800-722-1471 (TTY: 800-842-5357).

Український (Ukrainian):
Це повідомлення містить важливу інформацію. Це повідомлення може містити важливу інформацію про Ваше звернення щодо страхувального покриття через Premera Blue Cross. Зверніть увагу на ключові дати, які можуть бути вказані у цьому повідомленні. Існує імовірність того, що Вам треба буде здійснити певні кроки у конкретні кінцеві строки для того, щоб зберегти Ваше медичне страхування або отримати фінансову допомогу. У Вас є право на отримання цієї інформації та допомоги безкоштовно на Вашій рідній мові. Дозвоніться за номером телефону 800-722-1471 (TTY: 800-842-5357).

Tiếng Việt (Vietnamese):
Thông báo này cung cấp thông tin quan trọng. Thông báo này có thông tin quan trọng về đơn xin tham gia hoặc hợp đồng bảo hiểm của quý vị qua chương trình Premera Blue Cross. Xin xem kỹ các thông tin quan trọng trong thông báo này. Quý vị có thể phải thực hiện những bước sau trong thời hạn mà được dự trễ bảo hiểm sắp kết thúc hoặc đã được giúp đỡ từ trước khi thực hiện. Quý vị có quyền biết thông tin này và được truyền bá bằng ngôn ngữ của mình miễn phí. Xin gọi số 800-722-1471 (TTY: 800-842-5357).