



Coverage of any medical intervention discussed in a WellFirst Health medical policy is subject to the limitations and exclusions outlined in the member's benefit certificate or policy and to applicable state and/or federal laws.

Genetic Testing for Stickler Syndrome

MP9504

Covered Service: Yes

Prior Authorization Required: Yes

Additional Information: Genetic testing is covered for a WellFirst Health member if the test results provide a direct medical benefit or guides reproductive decision-making for the WellFirst Health member. See [Genetic Testing MP9012](#) for additional information.

Pre and post-test genetic counseling is required for any individual undergoing genetic testing.

For ASO members pre and post-genetic counseling is not required. Please reference the ASO Summary Plan Document (SPD).

A first-degree relative is defined as an individual's parents, full siblings, and children.

A second-degree relative is defined as an individual's grandparents, grandchildren, aunts, uncles, nephews, nieces and half-siblings.

WellFirst Health Medical Policy:

1.0 Stickler Syndrome sequencing panel – including COL2A1, COL11A1, COL11A2, COL9A1, COL9A2 and COL9A3 genetic testing **requires** prior authorization through the Health Services Division and is considered medically necessary when **2 or more** of the following are met:

- 1.1 Ocular findings – vitreous changes or retinal abnormalities (lattice degeneration, retinal hole, retinal detachment, or retinal tear)
- 1.2 High-frequency sensorineural hearing loss and/or frequent ear infections
- 1.3 Characteristic facial features including midfacial underdevelopment, malar hypoplasia, broad or flat nasal bridge, and micro/retrognathia
- 1.4 Cleft palate (open cleft, submucous cleft, or bifid uvula)

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- 1.5 Skeletal findings including:
 - 1.5.1 Osteoarthritis before age 40
 - 1.5.2 Slipped epiphysis or Legg-Perthes-like disease
 - 1.5.3 Scoliosis, spondylolisthesis, or Scheuermann-like kyphotic deformity
 - 1.5.4 First or second degree relative with a known pathogenic autosomal dominant or autosomal recessive sequence variant(s) in a related gene.
- 2.0 When criteria in 1.0 are met for testing of these genes we will cover deletion/duplication testing via array comparative genomic hybridization.
- 3.0 All other indications not listed above are considered experimental and investigational, and therefore are not covered.

CPT/HCPCS Codes Related to MP9504

The list of codes (and their descriptors, if any) is provided for informational purposes only and may not be all inclusive or current. Listing of a code in this medical policy does not imply that the service described by the code is a covered or non-covered service. Benefit coverage for any service is determined by the member's policy of health coverage with WellFirst Health. Inclusion of a code above does not imply any right to reimbursement or guarantee claim payment. Other medical policies may also apply.

CPT Code	Description
81479	Unlisted molecular pathology procedure



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