

	06/01/2024
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ADMINISTRA	ATIVE

Administrative Policy Statements prepared by CareSource and its affiliates do not ensure an authorization or payment of services. Please refer to the plan contract (often referred to as the Evidence of Coverage) for the service(s) referenced in the Administrative Policy Statement. If there is a conflict between the Administrative Policy Statement and the plan contract (i.e., Evidence of Coverage) will be the controlling document used to make the determination.

According to the rules of Mental Health Parity Addiction Equity Act (MHPAEA), coverage for the diagnosis and treatment of a behavioral health disorder will not be subject to any limitations that are less favorable than the limitations that apply to medical conditions as covered under this policy.

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The ADMINISTRATIVE Policy Statement detailed above has received due consideration as de



- IV. The following are not considered medically necessary and will not be covered:
 - A. repeat testing
 - B. gene sequencing of anything other than the CFTR gene
 - C. Fetal testing is generally not considered medically necessary. However, fetal testing is medically necessary with the presence of any of the following:
 - 1. Both parents have disease-causing mutations of the CFTR gene.
 - 2. Echogenic bowel is detected on ultrasound examination of fetus during pregnancy.
 - 3. The mother is a confirmed carrier, and father is unknown or unavailable for testing.
- E. Conditions of Coverage

CareSource may request documentation for post-payment review of claims submitted for payment of CF testing. If documentation is not provided, CareSource may recoup previously paid claim(s).

F. Related Policies/Rules Genetic Testing and Genetic Counseling

G. Review/Revision History

	DATES	ACTION
Date Issued	09/02/2020	
Date Revised	07/20/2022	Addition of Section D, IV and V.
	02/14/2024	Annual review: changes to title, background, and definitions, expanded policy to include diagnostic testing, and updated references. Approved at Committee.
Date Effective	06/01/2024	
Date Archived		

H. References

- Barben J, Castellani C, Munck A, et al. Updated guidance on the management of children with cystic fibrosis transmembrane conductance regulator-related metabolic syndrome/cystic fibrosis screen positive, inconclusive diagnosis (CRMS/CFSPID). J Cyst Fibros. 2021;20(5):810-819. doi:10.1016/j.jcf.2020.11.006
- Bienvenu T, Lopez M, Girodon E. Molecular diagnosis and genetic counseling of cystic fibrosis and related disorders: new challenges. *Genes (Basel)*. 2020;11(6):619. doi:10.3390/genes11060619
- Bombieri C, Claustres M, De Boeck K, et al. Recommendations for the classification of diseases as CFTR-related disorders. *J Cyst Fibros*. 2011;10(Suppl 2):S86-S102. doi:10.1016/S1569-1993(11)60014-3
- 4. Carrier. National Human Genome Research Institute. Updated October 15, 2023. Accessed December 28, 2023. www.genome.gov
- 5. Carrier testing for cystic fibrosis. Cystic Fibrosis Foundation. Accessed December 28, 2023. www.cff.org

The ADMINISTRATIVE Policy Statement detailed above has received due consideration as defined in the ADMINISTRATIVE Policy Statement Policy and is approved.