

Local Coverage Determination (LCD): MoIDX: Chromosome 1p/19q Deletion Analysis (L36483)

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Contractor Name	Contract Type	Contract Number	Jurisdiction	State(s)
Palmetto GBA	A and B and HHH	MAC 11201 - MAC A	J - M	South Carolina
Palmetto GBA	A and B and HHH	MAC 11202 - MAC B	J - M	South Carolina
Palmetto GBA	A and B and HHH	MAC 11301 - MAC A	J - M	Virginia
Palmetto GBA	A and B and HHH	MAC 11302 - MAC B	J - M	Virginia
Palmetto GBA	A and B and HHH	MAC 11401 - MAC A	J - M	West Virginia
Palmetto GBA	A and B and HHH	MAC 11402 - MAC B	J - M	West Virginia
Palmetto GBA	A and B and HHH	MAC 11501 - MAC A	J - M	North Carolina
Palmetto GBA	A and B and HHH	MAC 11502 - MAC B	J - M	North Carolina

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LCD Information

Document Information

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Original ICD-9 LCD ID
N/A

Revision Effective Date
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MoIDX: Chromosome 1p/19q Deletion Analysis

Revision Ending Date
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CMS National Coverage Policy CMS Internet Online Manual Pub. 100-02 (Medicare Benefit Policy Manual), Chapter 15, Section 80, "Requirements for Diagnostic X-Ray, Diagnostic Laboratory, and Other Diagnostic Tests"

CMS Internet-Only Manuals, Publication 100-04, Medicare Claims Processing Manual, Chapter 16, §50.5 Jurisdiction of Laboratory Claims, 60.12 Independent Laboratory Specimen Drawing, 60.2. Travel Allowance.

CMS Internet Online Manual Pub. 100-04 (Medicare Claims Processing Manual), Chapter 23 (Section 10) "Reporting ICD Diagnosis and Procedure Codes"

CMS Internet-Only Manual, Pub 100-04, Medicare Claims Processing Manual, Chapter 12, §30-Correct Coding Policy

Coverage Guidance

Coverage Indications, Limitations, and/or Medical Necessity

Indications for testing

Chromosome 1p-/19q- (eg, glial tumors), deletion analysis is considered medically necessary for the management of following glial tumors:

- Astrocytoma
- Ependymoma
- Oligoastrocytoma (Mixed Glioma)
- Oligodendroglioma
- Optic Glioma
- Gliomatosis Cerebri

Chromosome 1p-/19q-deletion analysis may also be indicated in the diagnosis of neoplasms that exhibit small round cell features (e.g. small glioblastomas and neurocytic tumors)

Limitations of coverage

Chromosome 1p-/19q- deletion analysis may be accomplished by molecular sequencing (81402) or morphometric analysis (e.g. in situ hybridization (FISH) 88367 or 88368). Physicians with patients who meet the indications of chromosome 1p-/19q testing - may select from one of the following test services:

- 81402 Chromosome 1p-/19q- (eg, glial tumors), deletion analysis
- 88367 Chromosome 1p-/19q- Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), using computer-assisted technology, per specimen; initial single probe stain procedure
- 8873 Chromosome 1p-/19q- Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), using computer-assisted technology, per specimen; each additional single probe stain procedure
- 88368 Chromosome 1p-/19q- Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), manual, per specimen; initial single probe stain procedure
- 88369 Chromosome 1p-/19q-Morphometric analysis, in situ hybridization (quantitative or semi-quantitative), manual, per specimen; each additional single probe stain procedure

Note: Only **ONE** chromosome 1p-/19q analysis service per patient will be considered reasonable and necessary

for tumor management.

Background

The presence of chromosome 1p/19q deletions in gliomas can assist in tumor differentiation, prognosis and treatment plan. Deletion studies specific to the 1p (short arm of chromosome #1) and 19q (long arm of chromosome #9) are performed on tumor tissue to determine if one or both 1p and 19q are deleted.

Over half of oligodendrogliomas have 1p/19q deletions that can help distinguish them from other types of gliomas.³ 1p/19q deletions can differentiate low-grade oligodendrogliomas from oligoastrocytomas.¹

The choice of adjuvant therapy depends on factors including tumor pathology and 1p/19q deletion status. Research observing improved survival has established combined procarbazine, lomustine, and vincristine (PCV) chemotherapy and radiation therapy as the new standard for treating anaplastic oligodendroglioma with the 1p/19q co-deletion.^{2,4,5,6}

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Coding Information

Bill Type Codes:

Contractors may specify Bill Types to help providers identify those Bill Types typically used to report this service. Absence of a Bill Type does not guarantee that the policy does not apply to that Bill Type. Complete absence of all Bill Types indicates that coverage is not influenced by Bill Type and the policy should be assumed to apply equally to all claims.

0x TBD

Revenue Codes:

Contractors may specify Revenue Codes to help providers identify those Revenue Codes typically used to report this service. In most instances Revenue Codes are purely advisory. Unless specified in the policy, services reported under other Revenue Codes are equally subject to this coverage determination. Complete absence of all Revenue Codes indicates that coverage is not influenced by Revenue Code and the policy should be assumed to apply equally to all Revenue Codes.

N/A

N/A

CPT/HCPCS Codes

Group 1 Paragraph: N/A

Group 1 Codes:

- 81402 MOLECULAR PATHOLOGY PROCEDURE, LEVEL 3 (EG, >10 SNPS, 2-10 METHYLATED VARIANTS, OR 2-10 SOMATIC VARIANTS [TYPICALLY USING NON-SEQUENCING TARGET VARIANT ANALYSIS], IMMUNOGLOBULIN AND T-CELL RECEPTOR GENE REARRANGMENTS, DUPLICATION/DELETION VARIANTS OF 1 EXON, LOSS OF HETEROZYGOSITY [LOH], UNIPARENTAL DISOMY [UPD])
- 88367 MORPHOMETRIC ANALYSIS, IN SITU HYBRIDIZATION (QUANTITATIVE OR SEMI-QUANTITATIVE), USING COMPUTER-ASSISTED TECHNOLOGY, PER SPECIMEN; INITIAL SINGLE PROBE STAIN PROCEDURE
- 88368 MORPHOMETRIC ANALYSIS, IN SITU HYBRIDIZATION (QUANTITATIVE OR SEMI-QUANTITATIVE), MANUAL, PER SPECIMEN; INITIAL SINGLE PROBE STAIN PROCEDURE
- 88369 MORPHOMETRIC ANALYSIS, IN SITU HYBRIDIZATION (QUANTITATIVE OR SEMI-QUANTITATIVE), MANUAL, PER SPECIMEN; EACH ADDITIONAL SINGLE PROBE STAIN PROCEDURE (LIST SEPARATELY IN ADDITION TO CODE FOR PRIMARY PROCEDURE)
- 88373 MORPHOMETRIC ANALYSIS, IN SITU HYBRIDIZATION (QUANTITATIVE OR SEMI-QUANTITATIVE), USING COMPUTER-ASSISTED TECHNOLOGY, PER SPECIMEN; EACH ADDITIONAL SINGLE PROBE STAIN PROCEDURE (LIST SEPARATELY IN ADDITION TO CODE FOR PRIMARY PROCEDURE)

ICD-10 Codes that Support Medical Necessity

Group 1 Paragraph: N/A

Group 1 Codes:

ICD-10 Codes	Description
C71.0	Malignant neoplasm of cerebrum, except lobes and ventricles
C71.1	Malignant neoplasm of frontal lobe
C71.2	Malignant neoplasm of temporal lobe
C71.3	Malignant neoplasm of parietal lobe
C71.4	Malignant neoplasm of occipital lobe
C71.5	Malignant neoplasm of cerebral ventricle
C71.6	Malignant neoplasm of cerebellum
C71.7	Malignant neoplasm of brain stem
C71.8	Malignant neoplasm of overlapping sites of brain
C71.9	Malignant neoplasm of brain, unspecified
C72.0	Malignant neoplasm of spinal cord

ICD-10 Codes that DO NOT Support Medical Necessity

Group 1 Paragraph: N/A

Group 1 Codes: N/A

ICD-10 Additional Information

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General Information

Associated Information

N/A

Sources of Information and Basis for Decision

1. Buckner JC, et al. Phase II trial of procarbazine, lomustine, and vincristine as initial therapy for patients with low-grade oligodendroglioma or oligoastrocytoma: efficacy and associations with chromosomal abnormalities. *J Clin Oncol.* 2003. 21(2):251-5.
2. Cairncross G, et al. Phase III trial of chemoradiotherapy for anaplastic oligodendroglioma: long-term results of RTOG 9402. *J Clin Oncol.* 2013. 31(3):337-43. doi: 10.1200/JCO.2012.43.2674. Epub.
3. Cairncross JG, et al. Specific genetic predictors of chemotherapeutic response and survival in patients with anaplastic oligodendrogliomas. *J Natl Cancer Inst.* 1998. 90(19):1473-9.
4. Hoang-Xuan K, et al. Temozolomide as initial treatment for adults with low-grade oligodendrogliomas or oligoastrocytomas and correlation with chromosome 1p deletions. *J Clin Oncol.* 2004. 22(15):3133-8.
5. Ino Y, et al. Molecular subtypes of anaplastic oligodendroglioma: implications for patient management at diagnosis. *Clin Cancer Res.* 2001. 7(4):839-45.
6. Kaloshi G, et al. Temozolomide for low-grade gliomas: predictive impact of 1p/19q loss on response and outcome. *Neurology.* 2007. 68(21):1831-6.

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Revision History Information

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Associated Documents

Attachments N/A

Related Local Coverage Documents Article(s) [A54776 - Response to Comments: MoIDX: Chromosome 1p/19q Deletion Analysis](#)

Related National Coverage Documents N/A

Public Version(s) Updated on 12/08/2015 with effective dates 02/01/2016 - N/A [Back to Top](#)

Keywords

N/A Read the [LCD Disclaimer](#) [Back to Top](#)