Local Coverage Determination (LCD): MoIDX: Molecular Diagnostic Tests (MDT) (L35160)

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Contractor Information

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
Noridian Healthcare Solutions, LLC	A and B MAC	01111 - MAC A	J - E	California - Entire State
Noridian Healthcare Solutions, LLC	A and B MAC	01112 - MAC B	J - E	California - Northern
Noridian Healthcare Solutions, LLC	A and B MAC	01182 - MAC B	J - E	California - Southern
Noridian Healthcare Solutions, LLC	A and B MAC	01211 - MAC A	J - E	American Samoa Guam Hawaii Northern Mariana Islands
Noridian Healthcare Solutions, LLC	A and B MAC	01212 - MAC B	J - E	American Samoa Guam Hawaii Northern Mariana Islands
Noridian Healthcare Solutions, LLC	A and B MAC	01311 - MAC A	J - E	Nevada
Noridian Healthcare Solutions, LLC	A and B MAC	01312 - MAC B	J - E	Nevada
Noridian Healthcare Solutions, LLC	A and B MAC	01911 - MAC A	J - E	American Samoa California - Entire State Guam Hawaii Nevada Northern Mariana Islands

LCD Information

Document Information

LCD ID

Original Effective Date

Original ICD-9 LCD ID

L33541

LCD Title

MoIDX: Molecular Diagnostic Tests (MDT)

Proposed LCD in Comment Period

N/A

Source Proposed LCD

N/A

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Revision Effective Date

For services performed on or after 01/01/2019

Revision Ending Date

N/A

Retirement Date

N/A

Notice Period Start Date

08/16/2015

Notice Period End Date

09/30/2015

CMS National Coverage Policy

Title XVIII of the Social Security Act (SSA) §1862(a)(1)(A), states that no Medicare payment shall be made for items or services that "are not reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of malformed body member."

Title XVIII of the Social Security Act (SSA) §1833(e), prohibits Medicare payment for any claim lacking the necessary documentation to process the claim.

Title XVIII of the Social Security Act (SSA) §1862(a)(1)(D), Investigational or Experimental.

CMS Manual System, Publication 100-02, *Medicare Benefit Policy Manual*, Chapter 15, §80.1, 80.1.1, 80.1.2, 80.1.3, laboratory services must meet applicable requirements of CLIA.

Pub 100-08 PIM, Ch. 13, Sec 13.1.3, Program Integrity Manual, "LCDs consist of only "reasonable and necessary" information.

Coverage Guidance

Coverage Indications, Limitations, and/or Medical Necessity

This coverage policy provides the following information:

- defines tests required to register for a unique identifier
- defines tests required to submit a complete technical assessment (TA) for coverage determination
- defines the payment rules applied to covered tests that are not reported with specific CPT codes
- lists some examples of specific covered tests that have completed the registration and TA process and meet Medicare's reasonable and necessary criteria for coverage. This listing is not inclusive.

Tests evaluated through the application process and/or technical assessment will be reviewed to answer the following questions:

- Is the test performed in the absence of clinical signs and symptoms of disease?
- Will the test results provide the clinician with information that will improve patient outcomes and/or change physician care and treatment of the patient?
- Will the test results confirm a diagnosis or known information?
- Is the test performed to determine risk for developing a disease or condition?
- Will risk assessment change management of the patient?
- Is there a diagnosis specific indication to perform the test?
- Is the test performed to measure the quality of a process or for Quality Control/Quality Assurance (QC/QA), i.e., a test to ensure a tissue specimen matches the patient?

MDT Policy Specific Definitions

MDT: Any test that involves the detection or identification of nucleic acid(s) (DNA/RNA), proteins, chromosomes, enzymes, cancer chemotherapy sensitivity and/or other metabolite(s). The test may or may not include multiple components. A MDT may consist of a single mutation analysis/identification, and/or may or may not rely upon an algorithm or other form of data evaluation/derivation.

LDT: Any test developed by a laboratory developed without FDA approval or clearance.

Applicable Tests/Assays

In addition to the MDT definition, this coverage policy applies to all tests that meet at least one of the following descriptions:

- All non-FDA approved/cleared laboratory developed tests (LDT)
- All modified FDA-approved/cleared kits/tests/assays
- All tests/assays billed with more than one CPT code to identify the service, including combinations of method-based, serology-based, and anatomic pathology codes
- All tests that meet the first three bullets and are billed with an NOC code

Unique Test Identifier Requirement

Because the available language in the HCPCS and CPT manuals to describe the pathology and laboratory categories and the tests included in those categories are not specific to the actual test results provided, all MDT services must include an identifier as additional claim documentation. Test providers must apply for an identifier specific to the applicable test and submit the test assigned identifier with the claim for reimbursement. The assigned identifier will provide a crosswalk between the test's associated detail information on file and the submitted claim detail line(s) required to adjudicate each test's claim. The unique identifier limits the need to submit the required additional information about the test on each claim.

Laboratory providers who bill MDT services must register services on the DEX™ Diagnostics Exchange.

<u>Technology Assessments (TA)</u>

MoIDX will review all new test/assay clinical information to determine if a test meets Medicare's reasonable and necessary requirement. Labs must submit a comprehensive dossier on each new test/assay prior to claim submission. MoIDX will only cover and reimburse tests that demonstrate analytical and clinical validity, and clinical utility at a level that meets the Medicare reasonable and necessary requirement.

Payment Rules

MoIDX will reimburse:

approved tests covered for dates of service consistent with the effective date of the coverage determination.

Covered Tests

Please refer to the Noridian website for specific coding and billing information for services addressed in MoIDX. .

Other tests/assays may be addressed by separate Noridian policy. In addition the CPT codes listed under Group 1 are addressed in the MoIDX program. If a test is not linked below under Related Local Coverage Documents, it may be addressed under separate Noridian policy or it has not been approved for coverage as it has either not been vetted by the MoIDx contractor or has been found to be considered statutorily excluded.

To obtain a unique identifier for a test and, to submit information for a technical assessment go to DEX™ Diagnostics Exchange:https://app.dexzcodes.com/login.

For additional MolDX Program information, go to the Noridian Medicare home page at noridianmedicare.com and select MolDX under the Policies Tab.

Coding Information			
NA			
(Rationale for Determination)			
Analysis of Evidence			
NA			
Summary of Evidence			
MoIDX expects laboratory providers to follow test	t indications published	by the developer.	

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.

N/A

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

CPT/HCPCS Codes

Group 1 Paragraph:

N/A

Group 1 Codes:

CODE	DESCRIPTION
81105 - 81112	Hpa-1 genotyping - Hpa-15 genotyping
81120 - 81121	Idh1 common variants - Idh2 common variants
81161 - 81599	Dmd dup/delet analysis - Unlisted maaa
84999	Clinical chemistry test
85999	Hematology procedure
86152 - 86153	Cell enumeration & id - Cell enumeration phys interp
86849	Immunology procedure
88120 - 88121	Cytp urne 3-5 probes ea spec - Cytp urine 3-5 probes cmptr
0002M	Liver dis 10 assays w/ash
0003M	Liver dis 10 assays w/nash
0004M	Scoliosis dna alys
0006M	Onc hep gene risk classifier
0007M	Onc gastro 51 gene nomogram
0009M	Fetal aneuploidy trisom risk
0011M	Onc prst8 ca mrna 12 gen alg
0012M	Onc mrna 5 gen rsk urthl ca
0013M	Onc mrna 5 gen recr urthl ca
0001U	Rbc dna hea 35 ag 11 bld grp
0002U	Onc clrct 3 ur metab alg plp
0003U	Onc ovar 5 prtn ser alg scor
0005U	Onco prst8 3 gene ur alg
0006U	Detc ia meds 120+ analytes
0007U	Rx test prsmv ur w/def conf
0008U	Hpylori detcj abx rstnc dna
0009U	Onc brst ca erbb2 amp/nonamp
0010U	Nfct ds strn typ whl gen seq
0011U	Rx mntr lc-ms/ms oral fluid
0012U	Germln do gene reargmt detcj
0013U	Onc sld org neo gene reargmt
0014U	Hem hmtlmf neo gene reargmt
0016U	Onc hmtlmf neo rna bcr/abl1
0017U	Onc hmtlmf neo jak2 mut dna

CODE	DESCRIPTION
0018U	Onc thyr 10 microrna seq alg
0019U	Onc rna tiss predict alg
0021U	Onc prst8 detcj 8 autoantb
0022U	Trgt gen seq dna&rna 23 gene
0023U	Onc aml dna detcj/nondetcj
0024U	Glyca nuc mr spectrsc quan
0025U	Tenofovir liq chrom ur quan
0026U	Onc thyr dna&mrna 112 genes
0027U	Jak2 gene trgt seq alys
0029U	Rx metab advrs trgt seq alys
0030U	Rx metab warf trgt seq alys
0031U	Cyp1a2 gene
0032U	Comt gene
0033U	Htr2a htr2c genes
0034U	Tpmt nudt15 genes
0035U	Neuro csf prion prtn qual
0036U	Xome tum & nml spec seq alys
0037U	Trgt gen seq dna 324 genes
0038U	Vitamin d srm microsamp quan
0039U	Dna antb 2strand hi avidity
0040U	Bcr/abl1 gene major bp quan
0041U	B brgdrferi antb 5 prtn igm
0042U	B brgdrferi antb 12 prtn igg
0043U	Tbrf b grp antb 4 prtn igm
0044U	Tbrf b grp antb 4 prtn igg

ICD-10 Codes that Support Medical Necessity

Group 1 Paragraph:

N/A

Group 1 Codes:

ICD-10 CODE	DESCRIPTION
XX000	Not Applicable

ICD-10 Codes that DO NOT Support Medical Necessity

N/A

Additional ICD-10 Information

N/A

General Information

Associated Information

N/A

Sources of Information

1. Current Procedural Terminology® (CPT) American Medical Association. American Medical Association Press, ISBN9781603592178, 2011.

Bibliography

NA

Revision History Information

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
01/01/2019	R9	The following paragraph under Covered Tests has been clarified: Other tests/assays may be addressed by separate Noridian policy. In addition the CPT codes listed under Group 1 are addressed in the MoIDX program. If a test is not linked below under Related Local Coverage Documents, it may be addressed under separate Noridian policy or it has not been approved for coverage as it has either not been vetted by the MoIDx contractor or has been found to be considered statutorily excluded.	Other (Clarification of paragraph requested by the MoIDX contractor.)
01/01/2019	R8	Corrected typographical error in R7 revision history: CPT codes 71178, 71179, 71180 should be 81178, 81179 and 81180.	Typographical Error
01/01/2019	R7	The following updates were made as a result of the 2019	Creation of Uniform LCDs With

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REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
		Annual HCPCS code update:	Other MAC Jurisdiction
		Deleted codes:0001M, 81211, 81213, 81214	 Revisions Due To CPT/HCPCS Code Changes
		Codes added to existing ranges: 81163, 81164, 81165, 81166, 81167, 81171, 81172, 81173, 81174, 81177, 71178, 71179, 71180, 81181, 81182, 81183, 81184, 81185, 81186, 81187, 81188, 81189, 81190, 81204, 81233, 81234, 81236, 81237, 81239, 81237, 81239, 81271, 81274, 81284, 81285, 81286, 81289, 81305, 81306, 81312, 80320, 81329, 81333, 81336, 81337, 81343, 81344,81345, 81443, 81518,81596	
		Codes with descriptor changes: 0006U, 0012M, 0031U, 0032U, 81109, 81162, 81212, 81215, 81216, 81217, 81244, 81287, 81327, 81334	
06/21/2018	R6	Removed: 88399, 89398, 87999, 88199, 88299	Creation of Uniform LCDs With
		Added: 0001U, 0002U, 0003U, 0005U, 0006U, 0007U, 0008U, 0009U, 0010U, 0011U, 0012U, 0013U, 0014U, 0016U, 0017U, 0018U, 0019U, 0020U, 0021U, 0022U, 0023U, 0024U, 0025U, 0026U, 0027U, 0028U, 0029U, 0030U, 0031U, 0032U, 0033U, 0034U, 0035U, 0036U, 0037U, 0038U, 0039U, 0040U, 0041U, 0042U, 0043U, 0044U, 0011M, 0012M, 0013M, 81105-81112, 81120-81121, 86152-86153, 88120-88121.	Other MAC Jurisdiction • Revisions Due To CPT/HCPCS Code Changes
01/01/2018	R5	Removed G0452, 88380, 88381 because they no longer require a DEX Z code identifier. Revised the link for technical assessment information.	 Creation of Uniform LCDs With Other MAC Jurisdiction
		03/29/2018: At this time 21st Century Cures Act will apply to new and revised LCDs that restrict coverage which requires comment and notice. This revision is not a restriction to the coverage determination; and, therefore not all the fields included on the LCD are applicable as noted in this policy.	
01/01/2018	R4	The following changes were made as a result of the Annual 2018 CPT/HCPCS code update:	 Creation of Uniform LCDs With Other MAC

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
		81175, 81176, 81230, 81231, 81232, 81238, 81247, 81248, 81249, 81258, 81259, 81269, 81283, 81328, 81334, 81335, 81346, 81361, 81362, 81363, 81364, 81448, 81520, 81521, 81541 and 81551 were added to code range 81161 - 81599 in Group 1.	Jurisdiction • Revisions Due To CPT/HCPCS Code Changes
		CPT codes are current as of the AMA CPT® 2018 Professional Edition, ISBN 978-1-62202-600-5, ISSN 0276-8283.	
		12/5/2017 At this time 21st Century Cures Act will apply to new and revised LCDs that restrict coverage which requires comment and notice. This revision is not a restriction to the coverage determination; and, therefore not all the fields included on the LCD are applicable as noted in this policy.	
01/01/2017	R3	2017 CPT Code Changes: The following CPT/HCPCS codes were added to these code ranges: 81327 was added to code range 81161 - 81599 in Group 1 81413 was added to code range 81161 - 81599 in Group 1 81414 was added to code range 81161 - 81599 in Group 1 81422 was added to code range 81161 - 81599 in Group 1 81439 was added to code range 81161 - 81599 in Group 1 81539 was added to code range 81161 - 81599 in Group 1 Description was changed for the following CPT/HCPCS codes: 81402 descriptor was changed in Group 1, 81407 descriptor was changed in Group 1 CPT/HCPCS codes were deleted:	Revisions Due To CPT/HCPCS Code Changes
04/21/2016	R2	0010M, 81280, 81281 and 81282 was deleted from Group 1. Replaced Palmetto GBA reference with MolDX, Under "Unique Test Identifier Requirement" - removed instruction to register services via Z-Code Identifier Application and Palmetto GBA Test Identifier (PTI) Application. Under "Payment Rules" - removed suspension of claims that omit Z-Code IDs. Under "Covered Tests" - updated the point of contact for McKesson and MolDX.) JEA LCD L36249 is retired and JEA contract numbers are added to the JEB LCD so that JEA and JEB have the same MCD LCD number.	Creation of Uniform LCDs With Other MAC Jurisdiction
10/01/2015	R1	This LCD is the final ICD-10 version of DL33541 Molecular Diagnostic Tests (MDT), which was initially introduced into draft in ICD-9 format and finalized in ICD-10 format.	Creation of Uniform LCDs

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
			Within a MAC Jurisdiction • Creation of Uniform LCDs With Other MAC Jurisdiction

Associated Documents

Attachments

N/A

Related Local Coverage Documents

Article(s)

A55711 - MolDX: Abbott RealTime IDH2 testing for Acute Myeloid Leukemia (AML) Billing and Coding Guidelines

A55881 - MolDX: ThermoFisher Oncomine Dx Target Test For Non-Small Cell Lung Cancer Billing and Coding

Guidelines

A54356 - MolDX: Afirma™ Assay by Veracyte Billing and Coding Guidelines

A54364 - MolDX: AlloMap Billing and Coding Guidelines

A54376 - MolDX: Avise PG Assay Billing and Coding Guidelines

A54386 - MolDX: bioTheranostics Cancer TYPE ID® Billing and Coding Guidelines

A55185 - MolDX: FDA Approved ALK Companion Diagnostic Tests Billing and Coding Guidelines

A54418 - MolDX: FDA-Approved BRAF Tests Billing and Coding Guidelines

A54422 - MolDX: FDA-Approved EGFR Tests Billing and Coding Guidelines

A54498 - MoIDX: FDA-Approved KRAS Tests

A54437 - MolDX: HERmark® Assay by Monogram Billing and Coding Guidelines

A54445 - MolDX: MammaPrint Billing and Coding Guidelines

A55294 - MolDX: Myriad's BRACAnalysis CDx™ Billing and Coding Guidelines

A54480 - MoIDX: Oncotype DX® Breast Cancer Assay Billing and Coding Guidelines

A54484 - MoIDX: Oncotype DX® Colon Cancer Coding and Billing Guidelines

A54489 - MolDX: Progensa® PCA3 Assay Billing and Coding Guidelines

A54494 - MolDx: ResponseDX Tissue of Origin® Billing and Coding Guidelines

A54503 - MolDX: Vectra™ DA Billing and Coding Guidelines

A54552 - Response to Comments: MolDX: Molecular Diagnostic Tests (MDT)

A54429

- (MCD Archive Site)A54509
- (MCD Archive Site)

Related National Coverage Documents

N/A

Public Version(s)

Updated on 01/23/2019 with effective dates 01/01/2019 - N/A

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Some older versions have been archived. Please visit the MCD Archive Site to retrieve them.

Keywords

- Afirma
- Allomap
- Avise PG
- Cancer TYPE ID
- cobas 4800 BRAF V600
- cobas EGFR
- ConfirmMDx Epigenetic Molecular Assay
- Corus CAD
- HERmark
- MammaPrint
- Oncotype DX Breast
- · Oncotype DX Colon
- Progensa PCA3
- therascreen EGFR
- therascreen KRAS
- · Tissue of Origin
- THXID BRAF V600E/K Test
- Vectra DA
- Vysis
- MoIDX