

Local Coverage Article: MoIDX: Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C) Testing Billing and Coding Guidelines (A54975)

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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 American Samoa
Noridian Healthcare Solutions, LLC	A and B MAC	01211 - MAC A	J - E	Guam Hawaii Northern Mariana Islands American Samoa
Noridian Healthcare Solutions, LLC	A and B MAC	01212 - MAC B	J - E	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 American Samoa California - Entire State
Noridian Healthcare Solutions, LLC	A and B MAC	01911 - MAC A	J - E	Guam Hawaii Nevada Northern Mariana Islands

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Article Information

General Information

Article ID A54975	Original Article Effective Date 06/01/2016
Article Title MoIDX: Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD/C) Testing Billing and Coding Guidelines	Revision Effective Date 06/01/2016
AMA CPT / ADA CDT / AHA NUBC Copyright Statement	Revision Ending Date N/A
	Retirement Date N/A

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Article Guidance

Article Text:

ARVD/C, characterized by fatty replacement of heart cells predominantly in the right ventricle of the heart, is most often inherited as an autosomal dominant disease that may be associated with testing in at least seven genes (RYR2, TMEM43, DSP, PKP2, DSG2, DSC2 and JUP). Genetic testing may be performed in panels of 5-7 of these genes and disease-causing mutation is expected to be identified in 42-55% of cases. Testing would be performed to confirm an established diagnosis or on individuals already diagnosed with ARVD/C to identify family members at risk. Therefore, MolDX has determined that testing for ARVD/C is a statutorily excluded test.

To receive an ARVD/C panel test denial, please submit the following claim information:

- An Advance Beneficiary Notice (ABN) is not required for statutorily excluded services
 - For a voluntary issued ABN, append with GX modifier
 - To indicate a statutorily excluded service, append with a GY modifier
- Select the appropriate diagnosis for the patient
- Enter the DEX Z-Code™ identifier adjacent to the CPT code in the comment/narrative field for the following Part B claim field/types:
 - Loop 2400 or SV101-7 for the 5010A1 837P
 - Item 19 for paper claim
- Enter the DEX Z-Code™ identifier adjacent to the CPT code in the comment/narrative field for the following Part A claim field/types:
 - Line SV202-7 for 837I electronic claim
 - Block 80 for the UB04 claim form

Reference: Sec. 1862 (1)(A) Statutory Exclusion covers diagnostic testing "except for items and services that are not reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member,..."

[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 article does not apply to that Bill Type. Complete absence of all Bill Types indicates that coverage is not influenced by Bill Type and the article 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 article, 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 article should be assumed to apply equally to all Revenue Codes.

N/A

CPT/HCPCS Codes

Group 1 Paragraph: N/A

Group 1 Codes:

Group 1 CPT/HCPCS Code	Group 1 CPT/HCPCS Code Description
81479	UNLISTED MOLECULAR PATHOLOGY PROCEDURE

ICD-10 Codes that are Covered N/A

ICD-10 Codes that are Not Covered N/A

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[Revision History Information](#)

Revision History Date Revision History Number Revision History Explanation

06/01/2016	R1	Added Part A billing instructions.
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[Back to Top](#) **Related Local Coverage Document(s)** LCD(s) [L36358 - MolDX: Biomarkers in Cardiovascular Risk Assessment](#)

Related National Coverage Document(s) N/A

Statutory Requirements URL(s) N/A

Rules and Regulations URL(s) N/A

CMS Manual Explanations URL(s) N/A

Other URL(s) N/A

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Keywords

- MoIDX
- Arrhythmogenic
- Ventricular
- Dysplasia
- Cardiomyopathy
- ARVD/C
- ARVDC
- 81479

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