

Local Coverage Article: MoIDX: ENG and ACVRL1 Gene Tests Coding and Billing Guidelines (A53536)

Links in PDF documents are not guaranteed to work. To follow a web link, please use the MCD Website.

Contractor Information

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
Palmetto GBA	A and B MAC	10111 - MAC A	J - J	Alabama
Palmetto GBA	A and B MAC	10112 - MAC B	J - J	Alabama
Palmetto GBA	A and B MAC	10211 - MAC A	J - J	Georgia
Palmetto GBA	A and B MAC	10212 - MAC B	J - J	Georgia
Palmetto GBA	A and B MAC	10311 - MAC A	J - J	Tennessee
Palmetto GBA	A and B MAC	10312 - MAC B	J - J	Tennessee
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

Article Information

General Information

Article ID

A53536

Original Effective Date

10/01/2015

Original ICD-9 Article ID[A53537](#)**Revision Effective Date**

01/01/2019

Article Title

MoIDX: ENG and ACVRL1 Gene Tests Coding and Billing Guidelines

Revision Ending Date

N/A

AMA CPT / ADA CDT / AHA NUBC Copyright Statement**Retirement Date**

N/A

CPT codes, descriptions and other data only are copyright 2018 American Medical Association. All Rights Reserved. Applicable FARS/HHSARS apply.

Current Dental Terminology © 2018 American Dental Association. All rights reserved.

Copyright © 2019, the American Hospital Association, Chicago, Illinois. Reproduced with permission. No portion of the AHA copyrighted materials contained within this publication may be copied without the express written consent of the AHA. AHA copyrighted materials including the UB-04 codes and descriptions may not be removed, copied, or utilized within any software, product, service, solution or derivative work without the written consent of the AHA. If an entity wishes to utilize any AHA materials, please contact the AHA at 312-893-6816. Making copies or utilizing the content of the UB-04 Manual, including the codes and/or descriptions, for internal purposes, resale and/or to be used in any product or publication; creating any modified or derivative work of the UB-04 Manual and/or codes and descriptions; and/or making any commercial use of UB-04 Manual or any portion thereof, including the codes and/or descriptions, is only authorized with an express license from the American Hospital Association. To license the electronic data file of UB-04 Data Specifications, contact Tim Carlson at (312) 893-6816 or Laryssa Marshall at (312) 893-6814. You may also contact us at ub04@healthforum.com.

Article Guidance

Article Text:

Hereditary hemorrhagic telangiectasia (HHT) is an autosomal dominant disorder with variable clinical presentation. Most common manifestations are epistaxis, recurrent gastrointestinal bleeding, iron deficiency anemia, and mucocutaneous telangectasias. Arteriovenous malformations (AVMs), of various organs (lung, liver, brain) are common.

The two most prevalent forms of HHT, Type 1 and Type 2, are caused by mutations in the endoglin (ENG) or the ACVRL1 gene respectively. Although identification of these gene mutations can confirm the diagnosis of HHT, these tests are not necessary in many cases. HHT is generally established using well vetted consensus criteria (most often the International Curaçao Criteria (ICC)). The ICC uses the clinical characteristics of epistaxis, cutaneous or mucosal telangectasias, visceral AVMs, and a first-degree relative with HHT to judge likelihood of a given patient having HHT. A 'definite' diagnosis is established when a patient has 3 or 4 of these criteria. Genetic testing for ENG/ACVRL1 is not warranted. A patient with 0 to 1 criteria is 'unlikely' to have HHT, and similarly would not be a candidate for genetic testing. Patients with 2 or 3 Curaçao criteria are defined as "suspected" of HHT and are candidates for ENG/ACVRL1

testing.

Since screening of patients without signs or symptoms of HHT, who have a first-degree relative with HHT, is not a Medicare benefit, the MoIDX Team has determined ENG and/or ACVRL1 genetic testing and panels of tests that include ENG/ACVRL1 are statutorily excluded services.

EXCEPTIONS: For patients with “suspected” HHT in which diagnosis confirmation would demonstrate an improved outcome, approval will be made on a case-by-case basis through the appeal process.

For tests that include ENG and ACVRL1 registered and assigned a single Z-Code™ Identifier, submit CPT code 81479.

To receive an ENG and/or ACVRL1 gene test denial, please submit the following claim information:

- Select appropriate CPT code for test
 - 81405-ENG, Duplication/deletion
 - 81406-ENG, full gene sequence
 - 81479-ACVRL1
 - 81479-ACVRL1 and ENG
- An Advance Beneficiary Notice (ABN) is not required for statutorily excluded services
 - For a voluntary issued ABN, append with GX modifier
 - To indicate a valid ABN is on file for a known statutorily excluded service, append with a GY modifier
- Select the appropriate diagnosis for the patient
- For CPT non-NOC codes, Labs may either use the SV101-7 or SV202-7 (preferred) or the NTE field to submit this required information.
 - Enter the appropriate 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
 - Box 19 for paper claim
 - Enter the appropriate 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 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

N/A

ICD-10 Codes that are Covered

N/A

ICD-10 Codes that are Not Covered

N/A

Revision History Information

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
01/01/2019	R7	Removed 81403 from the article. This is due to the 2019 CPT/HCPCS Annual update and is effective 1/1/19.
04/12/2018	R6	Conducted annual validation and corrected bullet issues. No change in article content.
02/26/2018	R5	The Jurisdiction "J" Part A and Part B Contracts for Alabama (10111/10112), Georgia (10211/10212) and Tennessee (10311/10312) are now being serviced by Palmetto GBA. Effective 02/26/18, these 6 contract numbers are being added to this article. No coverage, coding or other substantive changes (beyond the addition of the 6 Part A and B contract numbers) have been completed in this revision.
04/27/2017	R4	Annual review completed. Updated Part-A & Part B billing instructions.

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
11/19/2015	R3	Removed the "MoIDX ID (MID) field" and changed back to SV101-7 and replaced MoIDX ID with Z-Code™ Identifier
11/19/2015	R2	Replaced SV101-7 with MID, and removed Palmetto GBA reference
10/01/2015	R1	Updated Annual Review date.

Associated Documents

Related Local Coverage Document(s)

N/A

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

Public Version(s)

Updated on 12/21/2018 with effective dates 01/01/2019 - N/A

Updated on 04/05/2018 with effective dates 04/12/2018 - N/A

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

Keywords

N/A