



# Healthcare Common Procedure Coding System (HCPCS) Codes Subject to and Excluded from Clinical Laboratory Improvement Amendment (CLIA) Edits

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Note: We revised this article on March 24, 2020, to reflect an updated CR 11640. In the article, we revised the transmittal number, CR release date and link to the transmittal. All other information remains the same.

## PROVIDER TYPES AFFECTED

This MLN Matters Article is for physicians, providers and suppliers billing Medicare Administrative Contractors (MACs) for services they provide to Medicare beneficiaries.

### **PROVIDER ACTION NEEDED**

CR 11640 informs the MACs about new HCPCS codes for 2020 that are subject to and excluded from Clinical Laboratory Improvement Amendment (CLIA) edits. Please make sure your billing staffs are aware of this update.

### BACKGROUND

Note: On March 5, CMS released information about developing a second HCPCS billing code (U0002), which laboratories can use to bill for certain 2019-Novel Coronavirus (COVID-19) and SARS-Co-V-2 diagnostic tests to help increase testing and track new cases, in addition to a previous HCPCS billing code (U0001), which laboratories can use to test for SARS-Co-V-2. (See <a href="https://www.cms.gov/newsroom/press-releases/cms-develops-additional-code-coronavirus-lab-tests">https://www.cms.gov/newsroom/press-releases/cms-develops-additional-code-coronavirus-lab-tests</a>.)

CLIA regulations require facilities to be appropriately certified for each test performed. To ensure that Medicare and Medicaid only pay for laboratory tests performed in certified facilities, each claim for an HCPCS code that is considered a CLIA laboratory test is currently edited at the CLIA certificate level.



HCPCS codes that are considered a laboratory test under CLIA change annually.

The following HCPCS code was discontinued on June 30, 2019:

• 0057U – mRNA gene analysis of 51 genes in solid organ tumor tissue

The following HCPCS code was discontinued on September 30, 2019:

 0104U – Hereditary pan cancer (eg, hereditary breast and ovarian cancer, hereditary endometrial cancer, hereditary colorectal cancer), genomic sequence analysis panel utilizing a combination of ngs, sanger, mlpa, and array cgh, with mrna analytics to resolve variants of unknown significance when indicated (32 genes [sequencing and deletion/duplication], epcam and grem1 [deletion/duplication only])

The following HCPCS codes were discontinued on Decmeber 31, 2019:

- 0081U Oncology (uveal melanoma), mrna, gene-expression profiling by real-time rt-pcr of 15 genes (12 content and 3 housekeeping genes), utilizing fine needle aspirate or formalin-fixed paraffin-embedded tissue, algorithm reported as risk of metastasis
- 0085U Evaluation of biomarker antibodies for inflammatory bowel disease

The following HCPCS codes are new for 2020, are excluded from CLIA edits, and do not require a facility to have any CLIA certificate:

- 0106U Evaluation of gastric emptying by measurement of radiolabeled carbon monoxide in breath specimens
- 0126U Analysis of 5 substances in maternal blood to assess risk of preeclampsia
- 0128U Analysis of 3 substances in maternal blood and analysis of Y chromosome in fetal DNA to assess risk of abnormal chromosomes in fetus and preeclampsia
- 0156U Copy number (eg, intellectual disability, dysmorphology), sequence analysis

The HCPCS codes below were added on July 1, 2019, and are subject to CLIA edits. They require a facility to have either a CLIA certificate of registration (certificate type code 9), a CLIA certificate of compliance (certificate type code 1), or a CLIA certificate of accreditation (certificate type code 3). A facility without a valid and current CLIA certificate, with a current CLIA certificate of waiver (certificate type code 2), or with a current CLIA certificate for provider-performed microscopy procedures (certificate type code 4) is not permitted to be paid for these tests.

- 0087U mRNA gene expression profiling of genes in heart transplant biopsy tissue to evaluate risk of rejection
- 0088U mRNA gene expression profiling of genes in kidney transplant tissue to evaluate risk of rejection
- 0089U Gene expression profiling of melanoma in superficial sample collected by adhesive patch



- 0090U mRNA gene expression profiling of 23 genes in skin melanoma tissue sample
- 0092U Measurement of 3 protein biomarkers for lung cancer in plasma
- 0093U Prescription drug monitoring for 65 common drugs in urine
- 0094U Rapid sequence gene testing
- 0095U Test for markers of eosinophilic inflammation of esophagus
- 0096U Test for detection of high-risk human papillomavirus in male urine
- 0097U Test for detection of gastrointestinal disease-causing organism using amplified probe
- 0098U Test for detection of respiratory disease-causing organism using amplified probe, 14 target organisms
- 0099U Test for detection of respiratory disease-causing organism using amplified probe, 20 target organisms (adenovirus, coronavirus 229E, coronavirus HKU1, coronavirus, coronavirus OC43, human metapneumovirus, influenza A, influenza A subtype, influenza A subtype H3, influenza A subtype H1-2009, influenza, parainfluenza virus, parainfluenza virus 2, parainfluenza virus 3, parainfluenza virus 4, human rhinovirus/enterovirus, respiratory syncytial virus, Bordetella pertussis, Chlamydophila pneumonia, Mycoplasma pneumoniae)
- 0100U Test for detection of respiratory disease-causing organism using amplified probe, 20 target organisms (adenovirus, coronavirus 229E, coronavirus HKU1, coronavirus NL63, coronavirus OC43, human metapneumovirus, human rhinovirus/enterovirus, influenza A, including subtypes H1, H1-2009, and H3, influenza B, parainfluenza virus 1, parainfluenza virus 2, parainfluenza virus 3, parainfluenza virus 4, respiratory syncytial virus, Bordetella parapertussis [IS1001], Bordetella pertussis [ptxP], Chlamydia pneumoniae, Mycoplasma pneumoniae)
- 0101U Gene sequence analysis panel of 15 genes associated with hereditary colon cancer and related disorders
- 0102U Gene sequence analysis panel of 17 genes associated with hereditary breast cancer and related disorders
- 0103U Gene sequence analysis panel of 24 genes associated with hereditary ovarian cancer and related disorders

The HCPCS codes below were added on October 1, 2019, and are subject to CLIA edits. They require a facility to have either a CLIA certificate of registration (certificate type code 9), a CLIA certificate of compliance (certificate type code 1), or a CLIA certificate of accreditation (certificate type code 3). A facility without a valid and current CLIA certificate, with a current CLIA certificate of waiver (certificate type code 2), or with a current CLIA certificate for provider-performed microscopy procedures (certificate type code 4) is not permitted to be paid for these tests.

- 0105U Measurement of tumor necrosis factor receptor 1A, receptor superfamily 2 (TNFR1, TNFR2), and kidney injury molecule-1 (KIM-1) in plasma to evaluate risk of rapid kidney function decline
- 0107U Antigen test for detection of Clostridium difficile toxin in stool
- 0108U Computer-assisted digital imaging of esophagus specimen slides to evaluate risk of cancer



- 0109U DNA test for detection of 4 Aspergillus species
- 0110U Monitoring of anti-cancer drugs in patient blood, serum, or plasma
- 0111U Gene analysis (KRAS and NRAS) in prostate tumor tissue
- 0112U Gene analysis for detection of infectious agent and drug resistance gene
- 0113U Measurement of PCA3 gene in urine and prostate-specific antigen (PSA) in serum to evaluate risk of prostate cancer
- 0114U Gene analysis (VIM and CCNA1 methylation) in esophageal cells to evaluate likelihood of precancerous changes
- 0115U Respiratory infectious agent detection by nucleic acid (DNA and RNA), 18 viral types and subtypes and 2
- O116U Analysis of 35 or more drugs in mouth fluid to evaluate risk of prescription drug interactions
- 0117U Analysis of 11 biochemical substances in urine to evaluate likelihood of atypical biochemical function associated with pain
- 0118U Measurement of transplant donor cell-free DNA in transplant recipient plasma
- 0119U Measurement of ceramides for assessment of heart disease risk
- 0120U mRNA, gene expression profiling of 58 genes in tissue sample for B-cell lymphoma classification
- 0121U Blood test for sickle cells using VCAM-1
- 0122U Blood test for sickle cells using P-Selectin
- 0123U Test for fragility of red blood cells
- 0124U Analysis of 3 substances in maternal blood to assess risk of abnormal chromosomes in fetus
- 0125U Analysis of 5 substances in maternal blood to assess risk of abnormal chromosomes in fetus and preeclampsia
- 0127U Analysis of 3 substances in maternal blood to assess risk of preeclampsia
- 0129U Gene analysis of genes associated with hereditary breast cancer and related disorders for gene sequence and duplication or deletion variants
- 0130U Targeted mRNA sequence analysis of genes associated with hereditary colon cancer and related disorders
- 0131U Targeted mRNA sequence analysis of 13 genes associated with hereditary breast cancer and related disorders
- 0132U Targeted mRNA sequence analysis of 17 genes associated with hereditary ovarian cancer and related disorders
- 0133U Targeted mRNA sequence analysis of 11 genes associated with hereditary prostate cancer and related disorders
- 0134U Targeted mRNA sequence analysis of 18 genes associated with hereditary pan cancer
- 0135U Targeted mRNA sequence analysis of 12 genes associated with hereditary gynecological cancer
- 0136U mRNA gene analysis (ataxia telangiectasia mutated)
- 0137U mRNA gene analysis (partner and localizer of BRCA2)
- 0138U mRNA gene analysis (BRCA1, DNA repair associated and BRCA2, DNA repair associated)



The HCPCS codes below are new for 2020 and are subject to CLIA edits. This list does not include new HCPCS codes for waived tests or provider-performed procedures. These codes require a facility to have either a CLIA certificate of registration (certificate type code 9), a CLIA certificate of compliance (certificate type code 1), or a CLIA certificate of accreditation (certificate type code 3). A facility without a valid and current CLIA certificate, with a current CLIA certificate of waiver (certificate type code 2), or with a current CLIA certificate for provider-performed microscopy procedures (certificate type code 4) is not permitted to be paid for these tests.

- 0139U Neurology (autism spectrum disorder [asd]), quantitative measurements of 6 central carbon metabolites (ie, ketoglutarate, alanine, lactate, phenylalanine, pyruvate, and succinate), lc-ms/ms, plasma, algorithmic analysis with result reported as negative or positive abolic subtypes of asd)
- 0140U Infectious disease (fungi), fungal pathogen identification, dna (15 fungal targets), blood culture, amplified probe technique, each target reported as detected or not detected
- 0141U Infectious disease (bacteria and fungi), gram-positive organism identification and drug resistance element detection, dna (20 gram-positive bacterial targets, 4 resistance genes, 1 pan gram-negative bacterial target, 1 pan candida target), blood culture, amplified probe technique, each target reported as detected or not detected
- 0142U Infectious disease (bacteria and fungi), gram-negative bacterial identification and drug resistance element detection, dna (21 gram-negative bacterial targets, 6 resistance genes, 1 pan gram-positive bacterial target, 1 pan candida target), amplified probe technique, each target reported as detected or not detected
- 0143U Drug assay, definitive, 120 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0144U Drug assay, definitive, 160 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0145U Drug assay, definitive, 65 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0146U Drug assay, definitive, 80 or more drugs or metabolites, urine, by quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0147U Drug assay, definitive, 85 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple



reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service

- 0148U Drug assay, definitive, 100 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0149U Drug assay, definitive, 60 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0150U Drug assay, definitive, 120 or more drugs or metabolites, urine, quantitative liquid chromatography with tandem mass spectrometry (lc-ms/ms) using multiple reaction monitoring (mrm), with drug or metabolite description, comments including sample validation, per date of service
- 0151U Infectious disease (bacterial or viral respiratory tract infection), pathogen specific nucleic acid (dna or rna), 33 targets, real-time semi-quantitative pcr, bronchoalveolar lavage, sputum, or endotracheal aspirate, detection of 33 organismal and antibiotic resistance genes with limited semi-quantitative results
- 0152U Infectious disease (bacteria, fungi, parasites, and dna viruses), dna, pcr and next-generation sequencing, plasma, detection of ]1,000 potential microbialorganisms for significant positive pathogens
- 0153U Oncology (breast), mrna, gene expression profiling by next-generation sequencing of 101 genes, utilizing formalin-fixed paraffin-embedded tissue, algorithm reported as a triple negative breast cancer clinical subtype(s) with information on immune cell involvement
- 0154U Fgfr3 (fibroblast growth factor receptor 3) gene analysis (ie, p.r248c [c.742c]t], p.s249c [c.746c]g], p.g370c [c.1108g]t], p.y373c [c.1118a]g], fgfr3-tacc3v1, and fgfr3tacc3v3)
- 0155U Pik3ca (phosphatidylinositol-4,5bisphosphate 3-kinase, catalytic subunit alpha) (eg, breast cancer) gene analysis (ie, p.c420r, p.e542k, p.e545a, p.e545d [g.1635g]t only], p.e545g, p.e545k, p.q546e, p.q546r, p.h1047l, p.h1047r, p.h1047y)
- 0157U Apc (apc regulator of wnt signaling pathway) (eg, familial adenomatosis polyposis [fap]) mrna sequence analysis (list separately in addition to code for primary procedure)
- 0158U Mlh1 (mutl homolog 1) (eg, hereditary non-polyposis colorectal cancer, lynch syndrome) mrna sequence analysis (list separately in addition to code for primary procedure)
- 0159U Msh2 (muts homolog 2) (eg, hereditary colon cancer, lynch syndrome) mrna sequence analysis (list separately in addition to code for primary procedure)
- 0160U Msh6 (muts homolog 6) (eg, hereditary colon cancer, lynch syndrome) mrna sequence analysis (list separately in addition to code for primary procedure)
- 0161U Pms2 (pms1 homolog 2, mismatch repair system component) (eg, hereditary nonpolyposis colorectal cancer, lynch syndrome) mrna sequence analysis (list separately in addition to code for primary procedure)



- 0162U Hereditary colon cancer (lynch syndrome), targeted mrna sequence analysis panel (mlh1, msh2, msh6, pms2) (list separately in addition to code for primary procedure)
- 80145 Measurement of adalimumab
- 80187 Measurement of posaconazole
- 80230 Measurement of infliximab
- 80235 Measurement of lacosamide
- 80280 Measurement of vedolizumab
- 80285 Measurement of voriconazole
- 81277 Cancer cytogenomic array gene analysis
- 81307 Gene analysis (partner and localizer of BRCA2) full sequence analysis
- 81308 Gene analysis (partner and localizer of BRCA2) for detection of known familial variant
- 81309 Gene analysis (partner and localizer of BRCA2) targeted sequence analysis
- 81522 mRNA gene expression analysis of 12 genes in breast tumor tissue
- 81542 mRNA gene expression analysis of 22 genes in prostate tumor tissue
- 81552 mRNA gene expression analysis of 15 genes in eye melanoma tissue or fine needle aspirate
- 87563 Detection of Mycoplasma genitalium by DNA or RNA probe

In addition, for 2020, there was 1 new HCPCS code, 0091U (Colorectal cancer screening by enumeration of tumor cells in blood), that was not mentioned in Transmittal 4326, CR 11280, Quarterly Update for Clinical Laboratory Fee Schedule and Laboratory Services Subject to Reasonable Charge Payment. The testing described by this code is subject to the CLIA regulations; however, it is not payable by Medicare in CY 2020. Hence, this new code was not included in CR11640.

**Note**: MACs will not search their files to either retract payment for claims already paid or to retroactively pay claims. However, they will adjust claims that you bring to their attention.

#### ADDITIONAL INFORMATION

The official instruction, CR 11640, issued to your MAC regarding this change is available at <u>https://www.cms.gov/files/document/r10009cp.pdf</u>.

If you have questions, your MACs may have more information. Find their website at <u>http://go.cms.gov/MAC-website-list</u>.



### **DOCUMENT HISTORY**

Date of Change	Description
March 24, 2020	We revised this article to reflect a revised CR 11640. In the article, we revised the transmittal number, CR release date and link to the transmittal. All other information remains the same.
March 9, 2020	Initial article released.

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