

POLICY NUMBER	EFFECTIVE DATE	APPROVED BY
R20200008	8/01/2020	RPC (Reimbursement Policy Committee)

IMPORTANT NOTE ABOUT THIS REIMBURSEMENT POLICY:

ConnectiCare has policies in place that reflect billing or claims payment processes unique to our health plans. Current billing and claims payment policies apply to all our products, unless otherwise noted. ConnectiCare will inform you of new policies or changes in policies through updates to the Provider Manual and/or provider news. The information presented in this policy is accurate and current as of the date of this publication.

The information provided in ConnectiCare's policies is intended to serve only as a general reference resource for services described and is not intended to address every aspect of a reimbursement situation. Other factors affecting reimbursement may supplement, modify or, in some cases, supersede this policy. These factors may include, but are not limited to: legislative mandates, physician or other provider contracts, the member's benefit coverage documents and/or other reimbursement, medical or drug policies. Finally, this policy may not be implemented exactly the same way on the different electronic claims processing systems used by ConnectiCare due to programming or other constraints; however, ConnectiCare strives to minimize these variations.

ConnectiCare follows coding edits that are based on industry sources, including, but not limited to; CPT guidelines from the American Medical Association, specialty organizations, and CMS including NCCI and MUE. In coding scenarios where there appears to be conflicts between sources, we will apply the edits we determine are appropriate. ConnectiCare uses industry-standard claims editing software products when making decisions about appropriate claim editing practices. Upon request, we will provide an explanation of how ConnectiCare handles specific coding issues. If appropriate coding/billing guidelines or current reimbursement policies are not followed, ConnectiCare may deny the claim and/or recoup claim payment.

Overview

This policy addresses the ConnectiCare, Inc. reimbursement policies pertaining to clinical laboratory and related laboratory services (e.g., venipuncture and the handling and conveyance of the specimen to the laboratory) for provider claims submitted on a CMS-1500, whether performed in a physician's office, a hospital laboratory, or an independent laboratory. Note this policy does not address reimbursement for all laboratory codes. Coding relationships for laboratory topics not included within this policy are administered through ConnectiCare administrative and reimbursement policies. All services described in this policy may be subject to additional reimbursement policies.

If you are a physician, practitioner, or medical group, you may only bill for services that you or your staff perform. Pass-through billing is not permitted and may not be billed to our members. We only reimburse for laboratory services that you are certified to perform through the Federal Clinical Laboratory Improvement Amendments (CLIA). You must not bill our members for any laboratory services for which you lack the applicable CLIA certification. To validate whether a test requires CLIA visit CMS/FDA websites.



Policy statement:

Duplicate Laboratory Charges - Multiple Providers

Only one provider will be reimbursed when multiple providers bill identical services. ConnectiCare will reimburse the provider or entity that actually performed the test. Duplicate laboratory services are defined as identical or equivalent bundled laboratory codes.

Note: For the purpose of this policy, CPT codes 82947 and 82948 are not considered to be equivalent codes:

- 82947 Glucose; quantitative, blood (except reagent strip)
- 82948 Glucose; blood, reagent strip

Pathologist and Physician Laboratory Providers

If a pathologist and another physician or other qualified health care professional's offices submit identical laboratory codes for the same patient on the same date of service, only the pathologist's service is reimbursable.

Place of Service

The Place of Service (POS) identifies where the laboratory service was performed. ConnectiCare uses the codes indicated in the Centers for Medicare and Medicaid Services (CMS) Place of Service Codes for Professional Claims Database to determine if laboratory services are reimbursable.

Examples:

- If the physician bills for lab services performed in his/her office, the POS code 11 for "Office" is reported.
- If an independent laboratory bills for a test on a sample drawn on an inpatient or outpatient of a hospital, the POS code 81 for "Independent Laboratory" is reported.

Laboratory Panels

Individual laboratory codes, which together make up a laboratory panel code, will be combined into and reimbursed as the more comprehensive laboratory panel code as described under the specific laboratory panel headings below.

ConnectiCare also considers an individual component code included in the more comprehensive panel code when reported on the same date of service by the same individual physician or other qualified health care professional. The Professional Edition of the CPT book, Organ or Disease-Oriented Panel section states: "Do not report two or more panel codes that include any of the same constituent tests performed from the same patient collection. If a group of tests overlaps two or more panels, report the panel that incorporates the greater number of tests to fulfill the code definition and report the remaining tests using individual test codes."

In addition, it is not appropriate for a laboratory panel to be split amongst multiple laboratories or office/laboratory settings. This is also considered unbundling of a laboratory panel. Laboratory panels that have been split billed, or unbundled are not reimbursable.

Venipuncture and Specimen Collection

Specimen collection fees are not reimbursed when billed by the same provider who is rendering blood or related laboratory services.



Consistent with CMS, only one collection fee for each type of specimen per patient encounter, regardless of the number of specimens drawn, will be allowed. A collection fee will not be reimbursed to anyone who did not extract the specimen.

Venous blood collection by venipuncture and capillary blood specimen collection (CPT codes 36415 and 36416) will be reimbursed once per patient per date of service when reported by the Same Individual Physician or Other Qualified Health Care Professional. When CPT code 36416 is submitted with CPT code 36415, CPT code 36415 is the only venipuncture code considered eligible for reimbursement. No modifier overrides will exempt CPT code 36416 from bundling into CPT code 36415.

Consistent with CMS, ConnectiCare considers collection of a specimen from a completely implantable venous access device and from an established catheter (CPT codes 36591 and 36592) to be bundled into services assigned a CMS NPFS Status Indicator of A, R or T provided on the same date of service by the Same Individual Physician or Other Qualified Health Care Professional, for which payment is made. When CPT code 36591 is submitted with CPT code 36592, CPT code 36592 is the only venipuncture code considered eligible for reimbursement. No modifier overrides will exempt CPT code 36591 from bundling into CPT code 36592.

ConnectiCare considers venipuncture code S9529 (Routine venipuncture for collection of Specimen(s), single homebound, nursing home, or skilled nursing facility patient) a non-reimbursable service. The description for S9529 focuses on place of service for a service that is more precisely represented by CPT code 36415 and reported with the appropriate CMS place of service code.

Consistent with CMS, specimen collection HCPCS code G0471 is reimbursable only when a Specimen is collected from an individual in a skilled nursing facility or by a laboratory on behalf of a home health agency.

Laboratory Handling and Conveyance

CPT codes 99000 and 99001 and HCPCS codes H0048 and S3600 are included in the overall management of a patient and are not separately reimbursed when submitted with another code, or when submitted as the only code on a claim for the same date of service.

Code Q0091

HCPCS code Q0091 (screening Papanicolaou smear, obtaining, preparing, and conveyance of cervical or vaginal smear to laboratory) is eligible for reimbursement for Medicare beneficiaries only. For all other products it is considered to be part of the E/M and Pap smear codes and is not eligible for separate reimbursement.

Guidelines for Billing Units

When submitting multiple units of one code, the guidelines are based on code descriptions:

- If the CPT or HCPCS code description contains "per" or "each" or another unit of measurement and multiple services are provided, providers should bill the code on one line with the appropriate number of units.
- If the code does not contain a measurement such as "per" or "each" in the description of the code, providers should report one unit for all services.



- If a CPT or HCPCS code description does not contain "per" or "each," and multiples of that service are provided, providers may bill the code on one line with multiple units, or with the appropriate repeat service modifier on separate lines:
 - o The 76 modifier is used for repeat services
 - o The 91 modifier is used for clinical diagnostic laboratory tests

Modifiers

Technical/Professional Modifiers TC/26

- Technical/Professional Component Billing identifies proper coding of professional, technical and global procedures. Modifier 26 signifies the professional component of a procedure, and Modifier TC signifies the technical component.
- When the Centers for Medicare & Medicaid Services (CMS) National Physician Fee Schedule Relative Value File (NPFSRVF) designates that modifier 26 is applicable to a procedure code (PC/TC indicator of 1 or 6), and the procedure (e.g., laboratory) has been reported by a professional provider with a facility place of service, the procedure code must be reported with modifier 26 or it will not be eligible for reimbursement.
- When the NPFSRVF designates that the concept of a separate professional and technical component does not apply to a laboratory procedure (PC/TC indicator of 3 or 9), and a professional provider has reported the laboratory procedure code with a modifier 26 the laboratory procedure code will not be eligible for reimbursement. When a laboratory procedure with a PC/TC indicator of 3 or 9 is reported by a professional provider with a facility place of service, the laboratory procedure code will not be eligible for reimbursement since, in this case, the facility will bill for performing the laboratory procedure.
- A global laboratory procedure code includes reimbursement for both the professional and technical components.
 - When both components are performed by the same provider, the appropriate code must be reported without the 26/TC modifiers.
 - When a provider has reported a global procedure and also reported the same procedure with a professional (26) or technical component (TC) modifier on a different line or claim, the procedure reported with the 26 or TC modifier will not be eligible for reimbursement.
 - When a professional provider bills the global code (no modifiers) with a facility place of service, the code will not be eligible for reimbursement.
- CPT instructions state that modifier 59 should not be used when a more descriptive modifier is available. CMS guidelines cite that the -X {EPSU} modifiers are more selective versions of modifier 59 so it would be incorrect to include both modifiers on the same line. According to CMS and CPT coding guidelines, modifier 59, XE, XP, XS, or XU may be used when the same laboratory services are performed for the same patient on the same day. ConnectiCare will reimburse laboratory services reported with modifier 59, XE, XP, XS, or XU for different species or strains, as well as Specimens from distinctly separate anatomic sites.



Codes:

Separate Reimbursement will not be provided for the following services when performed by a nurse other ancillary staff:

CPT Codes	Description
36591	Collection of blood specimen from a completely implantable venous access device
36592	Collection of blood specimen using established central or peripheral catheter, venous, not otherwise specified

Separate Reimbursement will not be provided for the following:

CPT Codes	Description
99000	Handling and/or conveyance of specimen for transfer from the office to a laboratory
99001	Handling and/or conveyance of specimen for transfer from the patient in other than a physician's office to a laboratory (distance may be indicated)
99002	Handling, conveyance, and/or any other service in connection with the implementation of an order involving devices (eg, designing, fitting, packaging, handling, delivery or mailing) when devices such as orthotics, protectives, prosthetics are fabricated by an outside laboratory or shop but which items have been designed and are to be fitted and adjusted by the attending physician or other qualified health care professional
H0048	Alcohol and/or other drug testing: collection and handling only, specimens other than blood
Q0091	Screening Papanicolaou smear; obtaining, preparing and conveyance of cervical or vaginal smear to laboratory (eligible for reimbursement for Medicare beneficiaries only)
S3600	Stat laboratory request (situations other than S3601)

Reimbursement Guidelines for 87631-87633

Consistent with CMS guidelines, ConnectiCare will consider reimbursement for only 87631. ConnectiCare will not reimburse 87632-87633; any testing with 5+ targets (pathogens) is considered not medically necessary.

Respiratory virus testing is considered for reimbursement when submitted with a CPT or HCPCS code(s) other than 87632 or 87633.

CPT Codes	Description
87631	Infectious agent detection by nucleic acid (DNA or RNA); respiratory virus (eg, adenovirus, influenza virus, coronavirus, metapneumovirus, parainfluenza virus, respiratory syncytial virus, rhinovirus), includes multiplex reverse transcription, when performed, and multiplex amplified probe technique, multiple types or subtypes, 3-5 targets



CPT Codes	Description
87632	Infectious agent detection by nucleic acid (DNA or RNA); respiratory virus (eg, adenovirus, influenza virus, coronavirus, metapneumovirus, parainfluenza virus, respiratory syncytial virus, rhinovirus), includes multiplex reverse transcription, when performed, and multiplex amplified probe technique, multiple types or subtypes, 6-11 targets
87633	Infectious agent detection by nucleic acid (DNA or RNA); respiratory virus (eg, adenovirus, influenza virus, coronavirus, metapneumovirus, parainfluenza virus, respiratory syncytial virus, rhinovirus), includes multiplex reverse transcription, when performed, and multiplex amplified probe technique, multiple types or subtypes, 12-25 targets

CLIA (Clinical Laboratory Improvement Amendment) ID Requirement:

The Centers for Medicare & Medicaid Services (CMS) regulates all laboratory testing (except research) performed on humans in the U.S. through CLIA. Congress passed CLIA in 1988 to establish quality standards, strengthen Federal oversight of clinical laboratories, and ensure the accuracy and reliability of patient test results.

CLIA applies to all laboratories that examine "materials derived from the human body for the purpose of providing information for the diagnosis, prevention, or treatment of any disease or impairment of, or the assessment of the health of, human beings." (42 U.S.C. § 263a(a)).

CLIA mandates nearly all laboratories, including those in physician offices, must meet applicable Federal requirements and have a current CLIA certificate. CLIA applies to all entities providing clinical laboratory services including those that do not file Medicare test claims.

For purposes of this policy, a valid CLIA Certificate Identification number will be required for reimbursement of clinical laboratory services reported on a1500 Health Insurance Claim Form (a/k/a CMS-1500) or its electronic equivalent.

Any claim that does not contain the CLIA ID, invalid ID, and/or the complete servicing provider demographic information will be considered incomplete and rejected or denied. Claim line edits will also be applied if the lab certification level does not support the billed service code.

Laboratory service providers who do not meet the reporting requirements and/or do not have the appropriate level of CLIA certification for the services reported will not be reimbursed.

Additional information regarding CLIA, applying for or renewing a certificate, or regarding assigned test complexity levels can be found at the following website.

Clinical Laboratory Amendments (CLIA) Website

Reporting CLIA ID Number on a claim:

Report the billing laboratory's CLIA ID number in:

• Loop 2300, REF02, REF01=X4 for all CLIA-covered laboratory tests submitted on the claim

Laboratory service providers who do not meet the reporting requirements and/or do not have the appropriate level of CLIA certification for the services reported will <u>not</u> be reimbursed



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80158	80159	80161	80162	80163	80164	80165	80167	80168	80169	80170	80171
80173	80175	80176	80177	80179	80180	80181	80183	80184	80185	80186	80187
80188	80189	80190	80192	80193	80194	80195	80197	80198	80199	80200	80201
80202	80203	80204	80210	80230	80235	80280	80285	80299	80306	80307	80320
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85270	85280	85290	85291	85292	85293	85300	85301	85302	85303	85305	85306



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87496	87497	87498	87500	87501	87503	87505	87506	87507	87510	87511	87512
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87660	87661	87662	87797	87798	87799	87800	87802	87803	87810	87811	87850
87901	87902	87903	87904	87906	87910	87912	87999	88104	88106	88108	88112
88120	88121	88130	88140	88141	88142	88143	88147	88148	88150	88152	88153
88155	88160	88161	88162	88164	88165	88166	88167	88172	88173	88174	88175
88177	88182	88184	88185	88187	88188	88189	88199	88230	88233	88235	88237
88239	88245	88248	88249	88261	88262	88263	88264	88267	88269	88271	88272
88273	88274	88275	88280	88283	88285	88289	88291	88299	88300	88302	88304
88305	88307	88309	88311	88312	88313	88314	88319	88321	88323	88325	88331
88332	88333	88334	88341	88342	88344	88346	88348	88350	88355	88356	88358
88360	88361	88362	88363	88364	88365	88366	88367	88368	88369	88371	88372
88373	88374	88375	88377	88380	88381	88387	88388	88399	89050	89051	89060
89125	89160	89230	89240	89310	89320	89322	89325	89329	89330	89331	0001U
0002U	0003U	0005U	0007U	0008U	0009M	0009U	0010U	0011M	0011U	0012M	0012U
0013M	0013U	0014M	0014U	0015M	0016M	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	0045U	0046U	0047U
0048U	0049U	0050U	0051U	0052U	0053U	0054U	0055U	0056U	0058U	0059U	0060U
0062U	0063U	0064U	0065U	0067U	0068U	0069U	0070U	0071U	0072U	0073U	0074U
0075U	0076U	0077U	0078U	0080U	0082U	0083U	0105U	0107U	0108U	0109U	0110U
0111U	0112U	0113U	0114U	0115U	0116U	0117U	0118U	0119U	0120U	0121U	0122U
0123U	0129U	0130U	0131U	0132U	0133U	0134U	0135U	0136U	0137U	0138U	0139U
0140U	0141U	0142U	0143U	0144U	0145U	0146U	0147U	0148U	0149U	0150U	0151U
0152U	0153U	0154U	0155U	0157U	0158U	0159U	0160U	0161U	0162U	0163U	0164U
0165U	0166U	0167U	0168U	0169U	0170U	0171U	0172U	0173U	0174U	0175U	0176U
0177U	0178U	0179U	0180U	0181U	0182U	0183U	0184U	0185U	0186U	0187U	0188U
0189U	0190U	0191U	0192U	0193U	0194U	0195U	0196U	0197U	0198U	0199U	0200U
0201U	0203U	0204U	0205U	0206U	0207U	0208U	0209U	0210U	0211U	0212U	0213U
0214U	0215U	0216U	0217U	0218U	0219U	0220U	0221U	0222U	0227U	0228U	0229U
0230U	0231U	0232U	0233U	0234U	0235U	0236U	0237U	0238U	0239U	0420U	0421U
0422U	0423U	0424U	0425U	0426U	0427U	0428U	0429U	0430U	0431U	0432U	0433U
0434U	0435U	0436U	0437U	0438U	G0103	G0123	G0124	G0141	G0143	G0144	G0145
G0147	G0148	G0306	G0307	G0327	G0416	G0432	G0435	G0476	G0480	G0481	G0482
G0483	G0499	G0659	P3000	P3001	P7001		1				1



CLIA Waived Tests:

CLIA waived tests listed in the table below are laboratory testing/procedure codes that ConnectiCare will consider for reimbursement to its network physicians when performed in their office. This list represents the only laboratory testing/procedures that ConnectiCare network physicians may provide in their offices. All other laboratory testing/procedures must be performed by one of the participating laboratories in ConnectiCare's network.

Modifier QW must be appended to any CLIA waived tests when any applicable laboratory service is reported.

ConnectiCare will reimburse the CPT Codes listed below when performed in a physician's office. The CPT Codes below are for laboratory tests allowed with a CLIA Waiver (exempt from CLIA) and do not require a certificate of accreditation or compliance.

CLIA V	Vaived										
0240U	0241U	80047	80048	80051	80053	80061	80069	80178	80305	81000	81001
81002	81003	81007	81015	81020	81025	81050	81514	82010	82040	82043	82044
82120	82150	82247	82270	82271	82272	82274	82310	82330	82374	82435	82465
82523	82550	82565	82570	82679	82947	82950	82951	82952	82962	82977	82985
83001	83002	83026	83036	83037	83516	83605	83655	83718	83721	83861	83880
83986	84075	84132	84155	84295	84443	84450	84460	84478	84520	84550	84703
84830	85013	85014	85018	85025	85576	85610	85651	86294	86308	86318	86386
86580	86618	86701	86769	86780	86803	87077	87210	87338	87389	87400	87420
87430	87426	87428	87449	87502	87631	87634	87635	87636	87637	87651	87801
87804	87806	87807	87808	87809	87811	87880	87899	87905	88720	89220	89250
89251	89253	89254	89255	89257	89258	89259	89260	89261	89264	89268	89272
89280	89281	89291	89300	89321	89335	89337	89342	89343	89344	89346	89352
89353	89354	89356	92977	G0328	G0433	G0472	G0475**	U0002			

Note: **This code is acceptable for Medicare Plans only; please submit with appropriate CPT code for Commercial Plans

Laboratory Modifiers

Modifier	Description
90	 Reference (outside) laboratory. Modifier 90 indicates pass through billing for a service that was not performed by the billing provider. ConnectiCare will only reimburse providers for procedures that are performed by the same provider. ConnectiCare does not reimburse modifier 90



Modifier	Description
91	 Modifier 91 is appropriate when the repeat laboratory service is performed by a different individual in the same group with the same Federal Tax Identification number. According to CMS and CPT guidelines, Modifier 91 is appropriate when, during the course of treatment, it is necessary to repeat the same laboratory test for the same patient on the same day to obtain subsequent test results, such as when repeated blood tests are required at different intervals during the same day
92	 Alternative Laboratory Platform Testing. When laboratory testing is being performed using a kit or transportable instrument that wholly or in part consists of a single use, disposable analytical chamber, the service may be identified by adding modifier 92 to the usual laboratory procedure code (HIV testing 86701-86703, and 87389). The test does not require permanent dedicated space; hence by its design it may be hand carried or transported to the vicinity of the patient for immediate testing at that site, although location of the testing is not in itself determinative of the use of this modifier.
QW	CLIA Waived Test

Ordering MD Claim Requirements:

ConnectiCare may pend or deny your claim if you do not list the ordering provider. Diagnostic claims such as labs and/or radiology must include the ordering physician's name and NPI as well as TIN.

Definitions:

Term	Definition
CLIA (Clinical Laboratory Improvement Amendments)	The Centers for Medicare & Medicaid Services (CMS) regulates all laboratory testing (except research) performed on humans in the U.S. through the Clinical Laboratory Improvement Amendments (CLIA). In total, CLIA covers approximately 251,000 laboratory entities. The Division of Laboratory Services, within the Survey and Certification Group, under the Center for Clinical Standards and Quality (CCSQ) has the responsibility for implementing the CLIA Program. More information is available at: Clinical Laboratory Amendments (CLIA) Website
CLIA Waived Test	As defined by CLIA, waived tests are categorized as "simple laboratory examinations and procedures that have an insignificant risk of an erroneous result." https://www.cms.gov/Regulations-and-Guidance/Legislation/CLIA/Downloads/waivetbl.pdf
Venipuncture	Venipuncture is the process of withdrawing a sample of blood for the purpose of analysis or testing. There are several different methods for the collection of a blood sample. The most common method and site of venipuncture is the insertion of a needle into the cubital vein of the anterior forearm at the elbow fold.



Term	Definition
Multiple Component Blood Tests/Panels	The first entry in the Pathology and Laboratory Section of the Current Procedural Terminology (CPT®') codebook is labeled "Organ or Disease Oriented Panels." Under the code for each blood panel is an inclusive list of each component code which when grouped together comprise the entire blood panel. CPT indicates that these panels were developed for coding purposes only.
Duplicate Laboratory Service	Identical or equivalent bundled laboratory Component Codes, submitted for the same patient on the same date of service on separate claim lines or on different claims regardless of the assigned Maximum Frequency per Day (MFD) value.
Non-Reference Laboratory Provider	A physician reporting laboratory procedures performed in their office or a pathologist.
Physician Office Laboratory	A laboratory maintained by a physician or group of physicians for performing diagnostic tests in connection with the physician practice.
Independent Laboratory	An Independent Laboratory is one that is independent both of an attending or consulting physician's office and of a hospital that meets at least the requirements to qualify as an emergency hospital. An Independent Laboratory must meet Federal and State requirements for certification and proficiency testing under the Clinical Laboratories Improvement Act (CLIA). Independent Laboratory providers must append modifier 90 to all reported laboratory services.
Reference Laboratory	A Reference Laboratory that receives a Specimen from another, Referring Laboratory for testing and that actually performs the test is often referred to as an Independent Laboratory. Reference Laboratory providers must append modifier 90 to all reported laboratory services.
Referring Laboratory	A Referring Laboratory is one that receives a specimen to be tested and that refers the specimen to another laboratory for performance of the laboratory test. Referring Laboratory providers must append modifier 90 to all reported laboratory services.
Specimen	Tissue or tissues that is (are) submitted for individual and separate attention, requiring individual examination and pathological diagnosis. Two or more such Specimens from the same patient (eg, separately identifiable endoscopic biopsies, skin lesions) are each appropriately assigned an individual code reflective of its proper level of service.
Date of Service	The date of service (DOS) on a claim for a laboratory test is the date the Specimen was collected and if collected over 2 calendar days, the DOS is the date the collection ended.

References

- 1. https://www.cms.gov/medicare-coverage-database/view/article.aspx?articleid=58741&ver=9
- 2. https://www.cms.gov/files/document/mm11640.pdf
- 3. https://www.cms.gov/Regulations-and-Guidance/Legislation/CLIA/Downloads/SubjecttoCLIA.pdf
- 4. https://www.cms.gov/Regulations-and-Guidance/Legislation/CLIA/index
- 5. https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNProducts/Downloads/CLIABrochure.pdf
- 6. https://www.cms.gov/files/document/mm12131.pdf
- 7. https://www.cms.gov/files/document/mm12093.pdf



Revision history

DATE	REVISION
1/10/2024	Updated policy with the following, effective 1/1/2024:
	 Require CLIA Certification table: Added new codes 0420U, 0421U, 0422U, 0423U, 0424U, 0425U, 0426U, 0427U, 0428U, 0429U, 0430U, 0431U, 0432U, 0433U, 0434U, 0435U, 0436U, 0437U, 0438U, 81457, 81458, 81459, 81462, 81463, 81464, 81517, 82166, 86041, 86042, 86043, 86366, & 87523
	 Removed existing codes 81514, 86769, 87400, 87420, & 87430
	 CLIA Waived table: Added existing codes 81514, 86769, 87400, 87420, & 87430
10/24/2023	Updated to remove G9143 from policy
5/8/2023	 Require CLIA Certification Removed 87428 (effective 11/10/2020) CLIA Waived Added 87428 (effective 11/10/2020)
5/3/2023	 Removed deleted codes, effective 5/12/2023 Require CLIA Certification: 0224U & U0001
4/5/2023	 Removed deleted codes, effective 5/12/2023 Require CLIA Certification: U0001, U0003, U0004, & U0005
2/28/2023	 Reimbursement regarding 87631-87633 updated to align with CMS effective 7/14/2023.
	87632 and 87633 removed from Laboratory Tests Requiring CLIA Certification of compliance or accreditation table
	87631 moved to CLIA Waved table
2/14/2023	Removed non-covered codes 0202U, 0223U, and 0225U
12/2022	Updated policy to include "Reporting CLIA ID Number on a claim form"
10/2022	Updated policy to include: CLIA Waived: added CPT Code 87389 CLIA Waived table: added clarification that G0475 is acceptable for Medicare plans only; applicable CPT code must be billed for Commercial plan
11/2021	 Require CLIA Certification Removed 87801 (effective 1/01/2022) CLIA Waived Added 87801 (effective 1/01/2022)



DATE	REVISION
6/2021	 Require CLIA Certification Added G0327 (new code effective 7/01/2021) Removed 0204U & 02041U (effective 10/06/2020) CLIA Waived Added 0204U & 02041U (effective 10/06/2020)
2/2021	Removed deleted codes Require CLIA Certification: 81545 & 87450
1/2021	 In order to align with CMS updates, policy updated with the following: Removed deleted codes from list of codes requiring CLIA Certification: 0111T, 0006U, 0124U, 0125U & 0127U CLIA Waived: Added: 87636, 87637 & 87811 Require CLIA Certification: Added 80143,80151, 80161,80167, 80179,80181, 80189, 80193, 80204, 80210, 81168, 81191, 81192, 81193, 81194, 81278, 81279, 81338, 81339, 81347, 81348, 81351, 81352, 81353, 81357, 81360, 81419, 81513, 81514, 81529, 81546, 81554, 82077, 82681, 86328, 86408, 86409, 86413, 86769, 87428, 87811, 0014M, 0015M, 0016M, 0163U, 0164U, 0165U, 0166U, 0167U, 0168U, 0169U, 0170U, 0171U, 0172U, 0173U, 0174U, 0175U, 0176U, 0177U, 0178U, 0179U, 0180U, 0181U, 0182U, 0183U, 0184U, 0185U, 0186U, 0187U, 0188U, 0189U, 0190U, 0191U, 0192U, 0193U, 0194U, 0195U, 0196U, 0197U, 0198U, 0199U, 0200U, 0201U, 0204U, 0205U, 0206U, 0207U, 0208U, 0209U, 0210U, 0211U, 0212U, 0213U, 0214U, 0215U, 0216U, 0217U, 0218U, 0219U, 0220U, 0221U, 0222U, 0223U, 0224U, 0226U, 0227U, 0228U, 0229U, 0229U, 0230U, 0231U, 0232U, 0233U, 0234U, 0235U, 0236U, 0237U, 0238U, 0239U, 0240U, 0241U, U0001, U0003, U00004, U0005.
7/2020	 Updated policy with following COVID codes: CLIA Waived: U0002, 87426 & 87635 Require CLIA Certification: U0001, U0003, U0004, 0202U, 0223U, 0224U, 86328 & 86769
6/2020	 This policy replaces previous "In Office Laboratory Procedures" Updated policy to include CLIA ID Requirement Updated content to include CLIA Lab Codes; CLIA waived and those requiring CLIA certification Updated policy limiting CPT Codes 87631-87633 to facilities only Updated to include modifier 90 Updated to include Ordering MD Claim Requirement
5/2020	Reformatted and reorganized policy, transferred content to new template with new Reimbursement Policy Number