

# STAT outpatient laboratory services exceptions list

**Updated: Jan. 1, 2026**

## Preferred laboratory services protocol

According to the UnitedHealthcare Community Plan of Maryland Preferred Laboratory Services Protocol, laboratory services ordered for plan members by their primary care provider (PCP) or specialist must be performed at the outpatient medical laboratory designated on the member's health plan ID card. We developed this protocol to help our members access appropriate care and keep their health care costs down.

UnitedHealthcare will deny claims for services that aren't performed at the designated outpatient medical laboratory unless they qualify as an exception. Exceptions to the requirements include:

- Tests performed during a covered visit to an urgent care facility or hospital emergency department
- STAT tests performed during a covered visit to a care provider's office listed in the STAT Outpatient Laboratory Services Exception List (included on the next page). For purposes of this document, STAT refers to items that are urgent or emergent in nature.
- STAT tests necessary to perform services at the time of visit
- Pathology services performed on specimens obtained during surgery at a hospital outpatient department
- Tests required on an intraoperative or intra-procedure basis for outpatient surgery or outpatient procedures
- Preoperative blood type and crossmatch studies
- Situations in which services are preapproved and/or contract exceptions apply



## Questions?

Connect with us through chat 24/7 in the **UnitedHealthcare Provider Portal**.

## STAT laboratory tests

If laboratory results are required on a STAT basis, the designated outpatient medical laboratory can arrange quick pickup and reporting. If a care provider performs a STAT test for a UnitedHealthcare Community Plan member and bills for the service, they must use the ET modifier with the CPT® code for the test. Additionally, the diagnosis indicated on the claim must support the STAT billing.

The table on the following pages lists the STAT outpatient exceptions to our Preferred Laboratory Services Protocol with their corresponding CPT codes.

Exceptions to this policy include the following STAT tests:

CPT code	Description
0202U	Infectious disease (bacterial or viral respiratory tract infection), pathogen-specific nucleic acid (DNA or RNA), 22 targets, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), qualitative RT-PCR, nasopharyngeal swab, each pathogen reported as detected or not detected
0223U	Infectious disease (bacterial or viral respiratory tract infection), pathogen-specific nucleic acid (DNA or RNA), 22 targets, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), qualitative RT-PCR, nasopharyngeal swab, each pathogen reported as detected or not detected
0224U	Antibody, severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (Coronavirus disease [COVID-19]), includes titer(s), when performed (do not report 0224U in conjunction with 86769)
0225U	Infectious disease (bacterial or viral respiratory tract infection) pathogen-specific DNA and RNA, 21 targets, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), amplified probe technique, including multiplex reverse transcription for RNA targets, each analyte reported as detected or not detected
0226U	Surrogate viral neutralization test (sVNT), severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]), ELISA, plasma, serum
0240U	Infectious disease (viral respiratory tract infection), pathogen-specific RNA, 3 targets (severe acute respiratory syndrome coronavirus 2 [SARS-CoV-2], influenza A, influenza B), upper respiratory specimen, each pathogen reported as detected or not detected
0241U	Infectious disease (viral respiratory tract infection), pathogen-specific RNA, 4 targets (severe acute respiratory syndrome coronavirus 2 [SARS-CoV-2], influenza A, influenza B, respiratory syncytial virus [RSV]), upper respiratory specimen, each pathogen reported as detected or not detected
80048	Basic metabolic panel
80051	Electrolyte panel
80053	Comprehensive metabolic panel
80055	Obstetric panel
80061	Lipid panel

<b>CPT code</b>	<b>Description</b>
80069	Renal function panel
80074	Acute hepatitis panel
80076	Hepatic function panel
80081	Under organ or disease-oriented panels
80156	Carbamazepine drug assay
80162	Digoxin drug assay
80163	Therapeutic drug assays
80165	Therapeutic drug assays
80178	Lithium drug assay
80184	Phenobarbital drug assay
80185	Phenytoin: total drug assay
80195	Sirolimus
80198	Theophylline drug assay
81000	Urinalysis
81001	Urinalysis; automated w/microscopy
81002	Urinalysis; non-automated w/o microscopy
81003	Urinalysis; automated w/o microscopy
81005	Urinalysis; qualitative
81007	Urinalysis; bacteriuria screen, non-cultured
81015	Urinalysis microscopic only
81025	Urine pregnancy test; visual
82120	Amines, vaginal fluid
82150	Amylase
82247	Bilirubin; total
82248	Bilirubin; direct
82270	Blood occult; feces screening
82271	Blood occult; peroxidase activity qualitative, other
82272	Blood occult peroxidase activity qualitative, feces
82310	Calcium; total
82374	Carbon dioxide
82435	Chloride; blood
82465	Assay, blood/serum cholesterol
82550	Creatine kinase, total
82565	Creatinine, blood

<b>CPT code</b>	<b>Description</b>
<b>82800</b>	Gases; blood, pH only
<b>82803</b>	Gases; blood, pH, pCO <sub>2</sub> , pO <sub>2</sub> , CO <sub>2</sub> , HCO <sub>3</sub>
<b>82810</b>	Gases; blood O <sub>2</sub> saturation only
<b>82947</b>	Glucose; quantitative
<b>82948</b>	Glucose; blood reagent strip
<b>82962</b>	Glucose; blood by glucose monitoring device
<b>83516</b>	Immunoassay for analyte other than infectious agent antibody or infectious agent antigen; qualitative or semiquantitative, multiple-step method
<b>83615</b>	Lactate dehydrogenase (LD), (LDH)
<b>83718</b>	Assay of lipoprotein
<b>83735</b>	Magnesium
<b>83861</b>	Microfluidic analysis utilizing an integrated collection and analysis device, tear osmolarity
<b>84030</b>	Phenylalanine (PKU); blood
<b>84075</b>	Phosphatase; alkaline
<b>84100</b>	Phosphorus inorganic (phosphate)
<b>84132</b>	Potassium; serum, plasma or whole blood
<b>84295</b>	Sodium; serum plasma or whole blood
<b>84450</b>	Transferase; aspartate amino
<b>84460</b>	Transferase; alanine amino
<b>84520</b>	Urea nitrogen; quantitative
<b>84545</b>	Urea nitrogen; clearance
<b>84550</b>	Uric acid; blood
<b>84702</b>	Gonadotropin, chorionic; quantitative
<b>84703</b>	Gonadotropin, chorionic; qualitative
<b>85014</b>	Blood count: hematocrit (Hct)
<b>85018</b>	Blood count; hemoglobin (Hgb)
<b>85025</b>	Blood count; complete (CBC) hemogram and platelet
<b>85027</b>	Blood count; complete (CBC), automated with platelet count
<b>85049</b>	Hematology and coagulation
<b>85060</b>	Blood smear, peripheral, interpretation by physician
<b>85610</b>	Prothrombin time
<b>85730</b>	Thromboplastin time; partial (PTT)
<b>85732</b>	Thromboplastin time; partial (PTT), substitution
<b>86308</b>	Heterophile antibodies; screening

<b>CPT code</b>	<b>Description</b>
<b>86318</b>	Immunoassay infectious agent antibody
<b>86328</b>	Immunoassay for infectious agent antibody(ies), qualitative or semiquantitative, single step method (e.g., re-agent strip); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19])
<b>86408</b>	Neutralizing antibody, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]); screen
<b>86409</b>	Neutralizing antibody, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]); titer
<b>86413</b>	Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]) antibody, quantitative
<b>86580</b>	Skin test; tuberculosis, intradermal
<b>86703</b>	Antibody; HIV-1&HIV-2 SINGLE
<b>86769</b>	Antibody; severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19])
<b>86901</b>	Blood typing, serologic; Rh (D)
<b>87205</b>	Smear; gram or giemsa routine stain
<b>87210</b>	Smear; wet mount for infectious agents
<b>87220</b>	Tissue examination by KOH slide
<b>87400</b>	Infectious agent antigen detection by immunoassay technique, (e.g., enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], fluorescence immunoassay [FIA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative; influenza, A or B, each
<b>87420</b>	Infectious agent antigen detection by immunoassay technique, (e.g., enzyme immunoassay, enzyme-linked immunosorbent assay, immunochemiluminometric assay) qualitative or semi-quantitative, multiple-step method; respiratory syncytial virus
<b>87426</b>	Infectious agent antigen detection by immunoassay technique, (e.g., enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative, multiple-step qualitative or semiquantitative, multiple-step method; severe acute respiratory syndrome coronavirus (e.g., SARS-CoV, SARSCoV-2 coronavirus) (e.g., SARS-CoV, SARSCoV-2 [COVID-19])
<b>87428</b>	Severe acute respiratory syndrome coronavirus (e.g., SARS-CoV, SARS-CoV-2 [COVID19]) and influenza virus types A and B
<b>87430</b>	Infectious agent antigen detection by immunoassay technique, (e.g., enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], fluorescence immunoassay [FIA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative; Streptococcus, group A

<b>CPT code</b>	<b>Description</b>
<b>87635</b>	Infectious agent detection by nucleic acid (DNA or RNA); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]), amplified probe technique
<b>87636</b>	Severe acute respiratory syndrome coronavirus 2 (SARS-CoV2) (coronavirus disease [COVID-19]) antibody, quantitative
<b>87637</b>	Infectious agent detection by nucleic acid (DNA or RNA); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]), influenza virus types A and B, and respiratory syncytial virus, multiplex amplified probe technique
<b>87807</b>	Infectious agent antigen detection by immunoassay with direct optical observation; respiratory syncytial virus
<b>87809</b>	Infectious agent antigen detection by immunoassay with direct optical observation; adenovirus
<b>87811</b>	Infectious agent antigen detection by immunoassay with direct optical (e.g., immunoassay with direct optical (e.g., visual) observation); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19])
<b>88172</b>	Cytopathology; evaluation of fine needle aspirate
<b>88173</b>	Cytopathology, needle aspirate, cytology interpretation
<b>88738</b>	Pathology and laboratory/in vivo (transcutaneous) lab
<b>89300</b>	Semen analysis w/Huhner's test (post coital)
<b>83655QW*</b>	Capillary lead
<b>84703QW*</b>	Rapid pregnancy
<b>87502QW*</b>	Point of Care Molecular Influenza A&B
<b>87634QW*</b>	Point of Care RSV
<b>87804QW*</b>	Rapid influenza
<b>87880QW*</b>	Rapid streptococcus, group A
<b>86803QW*</b>	Hepatitis C antibody
<b>G2023</b>	Specimen collection for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]), any specimen source
<b>G2024</b>	Specimen collection for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]), from an individual in an SNF (skilled nursing facility) or by a laboratory on behalf of an HHA (home health agency), any specimen source
<b>Q0111</b>	Wet mounts including preparation of specimens
<b>Q0112</b>	All potassium hydroxide (KOH)
<b>U0001</b>	CDC 2019 novel coronavirus (2019-nCoV) real time RT-PCR diagnostic panel

CPT code	Description
U0002	2019-nCoV coronavirus, SARS-CoV-2/2019-nCoV (COVID-19), any technique, multiple types, or subtypes (includes all targets), non-CDC
U0003**	Infectious agent detection by nucleic acid (DNA or RNA); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]), amplified probe technique, making use of high throughput technologies as described by CMS-2020-01-R
U0004**	2019-nCoV coronavirus, SARS-CoV-2/2019-nCoV (COVID-19), any technique, multiple types or subtypes (includes all targets), non-CDC, making use of high-throughput technologies as described by CMS-2020-01-R

\* These tests use a methodology of the Clinical Laboratory Improvement Amendments (CLIA)-waived category and may sometimes be billed with the QW modifier. Excludes places of service 11, 21 and 22 for pathology.

\*\* When billing for Healthcare Common Procedure Coding System (HCPCS) codes U0003 and U0004, please note the following:

- U0003 should identify tests performed with high-throughput technologies that would otherwise be billed using CPT code 87635
- U0004 should identify tests performed with high-throughput technologies that would otherwise be billed using U0002
- Neither U0003 nor U0004 should be billed for tests that detect COVID-19 antibodies. When billing all codes listed above, an ET modifier is required.

CPT® is a registered trademark of the American Medical Association.