
This article replaces the September 2021 guidance. Updates are highlighted in yellow.

The services in this guidance document are currently reimbursable by NYS Medicaid fee-for-service (FFS) and Medicaid Managed Care (MMC) Plans. The fees* below are specific to Medicaid fee-for-service. For individuals enrolled in Medicaid Managed Care, providers should check with the individual’s plan for implementation details and billing guidance. Providers are reminded COVID-19 tests performed must be Food and Drug Administration (FDA) approved or granted Emergency Use Authorization (EUA) through the FDA and in agreement with the level of complexity assigned by Wadsworth Lab. Coverage of COVID-19 tests for diagnostic and screening (including administration) must be consistent with the recommendations of the Centers of Disease Control and Prevention (CDC).

* The fees and effect dates below are current as of December 2021. Providers should periodically check their respective fee schedules in eMedNY for updates: https://www.emedny.org/ProviderManuals/index.aspx

Complexity levels are available at the following link: https://www.cdc.gov/clia/test-complexities.html


Please note:

COVID-19 test codes not outlined in this guidance are not covered.

Persons with signs or symptoms of COVID-19 should have diagnostic testing.

The CDC also recommends testing for the following individuals: https://www.cdc.gov/coronavirus/2019-ncov/hcp/testing-overview.html

Providers are reminded CDC guidance states: "All persons being tested, regardless of results, should receive counseling on the continuation of risk reduction behaviors that help prevent the transmission of SARS-CoV-2 (e.g., wearing masks, physical distancing, avoiding crowds and poorly ventilated spaces)."

Providers who are already receiving payment from another source for COVID-19 testing, specimen collection, or monoclonal antibody infusion are not eligible for reimbursement from Medicaid for those tests, specimen collections, or infusions.

Reminder

Providers are prohibited from charging Medicaid members a co-payment or any cost sharing responsibility for specimen collection or testing to diagnose or screen for COVID-19, or for monoclonal antibody infusions to treat a SARS-CoV-2 infection. Providers are reminded to check the emergency indicator box on claims submissions to waive cost sharing for these services.
**Home-based testing:**
COVID-19 diagnostic tests with "at-home" sample collection are eligible for reimbursement when the criteria outlined in this guidance are met and the test is processed in a New York State-approved laboratory. Additionally, over-the-counter (OTC) FDA-Approved COVID-19 at-home test kits that are ordered by authorized Medicaid providers are eligible for reimbursement. For details on coverage of point-of-care tests with at-home results, please see Medicaid pharmacy guidance at the following link: [https://www.health.ny.gov/health_care/medicaid/covid19/docs/guidance_home_covid_testing.pdf](https://www.health.ny.gov/health_care/medicaid/covid19/docs/guidance_home_covid_testing.pdf).

**Molecular/PCR Tests:**

- **87635** (effective 3/13/2020) – 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.  
  \[FFS \text{ fee } = $51.31\]

- **U0002** (effective 3/13/2020) – 2019-nCoV Coronavirus, SARS-CoV-2/2019-nCov (COVID-19), any technique, multiple types or subtypes (includes all targets), non-CDC.  
  \[FFS \text{ fee } = $51.31\]

**High Throughput Tests:**

These tests utilize highly sophisticated throughput machines which require more intensive technician training (to ensure the role of extremely skilled personnel) and more time intensive processes (to assure quality). A high throughput technology uses a platform that employs automated processing of more than two hundred specimens a day.

It is noted that U0003 should identify tests that would otherwise be identified by CPT code 87635 but for being performed with these high throughput technologies. It is further noted that U0004 should identify tests that would otherwise be identified by U0002 but for being performed with these high throughput technologies. Finally, it is noted that neither U0003 nor U0004 should be used for tests that detect COVID-19 antibodies. ([https://www.cms.gov/files/document/cms-2020-01-r.pdf](https://www.cms.gov/files/document/cms-2020-01-r.pdf))

- **U0003** (effective 4/14/2020) – 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.  
  \[FFS \text{ fee } = $100 \text{ (until 12/31/2020), } $75 \text{ (as of 01/01/2021)}\]

- **U0004** (effective 4/14/2020) – 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.  
  \[FFS \text{ fee } = $100 \text{ (until 12/31/2020), } $75 \text{ (as of 01/01/2021)}\]

In accordance with CMS, the fees for high throughput tests were reduced to $75.
effective 01/01/2021. For dates of service on or after 01/01/2021, U0005 may be billed as an add on code, when appropriate.

- **U0005** (effective 01/01/2021) – 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, CDC or non-CDC, making use of high throughput technologies, completed within 2 calendar days from date of specimen collection (List separately in addition to either HCPCS code U0003 or U0004) as described by CMS-2020-01-R2.
  FFS fee = $25

**Antigen Tests:**

- **87426** (effective 06/25/2020) – 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 method; severe acute respiratory syndrome coronavirus (e.g., SARS-CoV-2, SARS-CoV-2 [COVID-19]).
  FFS fee = $45.23

- **87811** (effective 10/06/2020) – Infectious agent antigen detection by immunoassay with direct optical (i.e., visual) observation; severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]).
  FFS fee = $41.38

**Multiplex Tests:**

- **87428** (effective 11/10/2020) – 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; severe acute respiratory syndrome coronavirus (e.g., SARS-CoV-2, SARS-CoV-2 [COVID-19]) and influenza virus types a and b.
  FFS fee = $73.49

- **87636** (effective 10/06/2020) – Infectious agent detection by nucleic acid (DNA or RNA); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]) and influenza virus types a and b, multiplex amplified probe technique.
  FFS fee = $142.63

- **87637** (effective 12/01/2021) – 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.
  FFS fee = $142.63
Antibody Tests:

Please see the following link for information on antibody testing for NYS residents:

- **86328** (effective 4/10/2020) – Immunoassay for infectious agent antibody(ies), qualitative or semiquantitative, single step method (e.g., reagent strip); severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]).
  
  *FFS fee = $45.23*

- **86769** (effective 4/10/2020) – Antibody; severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]).
  
  *FFS fee = $42.13*

**Specimen Collection** (effective 05/22/2020): During the period of the emergency separate Medicaid reimbursement is available for specimen collection when this is the only service being performed. Providers billing for reimbursement of one of the above tests should not bill separately for specimen collection or report. These specimen collection components are included in reimbursement for the test. Providers/Clinics billing for other primary procedures for the same patient on the same day should not bill for specimen collection. For more information, please see the chart below.

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
<th>Practitioner/ Clinic (Non-FQHC) Reimbursement</th>
<th>FQHC Reimbursement</th>
</tr>
</thead>
</table>
| G2023| Specimen collection for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]) | $23.46                                        | Physician/PA/NP/Midwife: bill rate code 4012 when **specimen collection only** is provided. Offsite visit rate ($64.97 upstate/$72.73 downstate) will be paid.  
Physician/PA/NP/Midwife: bill rate code 4013 **when specimen collection and E&M are provided**. Full PPS rate will be paid.  
RN/LPN: bill Procedure Code G2023 (ordered ambulatory) **for specimen collection only**. A fee of $23.46 will be paid.  
All FQHC services in this chart are eligible for wrap payments |

Clinics should bill the codes outlined in this guidance via the ordered
ambulatory fee schedule. The COVID-19 test and specimen collection codes are not payable under ambulatory patient groups (APGs).

**Skilled Nursing Facilities** services are paid at a daily rate; therefore, separate reimbursement is not available.

**Certified Home Health Agencies (CHHAs)** should perform specimen collection as part of nursing visits when ordered for existing clients who receive nursing services.

**CHHA specimen collection for homebound patients who do not receive nursing services** is eligible for reimbursement on or after 11/01/2020. Please see the chart below for details.

<table>
<thead>
<tr>
<th>Rate Codes Used for CHHAs Only</th>
<th>Descriptions</th>
<th>CHHA Patient Receiving Nursing Services</th>
<th>CHHA Patient Receiving Personal Care Services</th>
<th>NON-CHHA Homebound Patient</th>
<th>MLTC Patient Homebound Patient</th>
</tr>
</thead>
<tbody>
<tr>
<td>4921</td>
<td>Specimen collection for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]), from a homebound*** individual on behalf of a home health agency, any specimen source(s). Reimburses $25.46. For CHHA use only.</td>
<td>X</td>
<td>✔</td>
<td>✔</td>
<td>✔</td>
</tr>
</tbody>
</table>

Do not bill for specimen collection or travel. This should be done as part of a nurse visit and is included in the CHHA rate.

CHHAs providing nursing staff to collect a specimen from their patient who only receives lower-level services may bill rate code 4921 and rate code 4922 (travel each way) if specimen collection is the **only** service performed.

CHHAs providing specimen collection to a homebound individual who is not a CHHA patient may bill rate code 4921 and rate code 4922 (travel each way) if specimen collection is the **only** service performed.

CHHAs providing specimen collection for homebound patients may bill rate code 4921 and rate code 4922 (travel each way) if specimen collection is the **only** service performed.
Rate Codes Used for CHHAs Only | Descriptions | CHHA Patient Receiving Nursing Services | CHHA Patient Receiving Personal Care Services | NON-CHHA Homebound Patient | MLTC Patient Homebound Patient
--- | --- | --- | --- | --- | ---
4922 | Travel for COVID-19 specimen collection. Reimburses $9.35 each way. For CHHA use only. When seeing multiple patients in the same location**, only bill 1 trip charge for the first Medicaid member visit. |  |  |  |  

** The same location is defined as a vehicle is not necessary to travel between visits.  
*** Patients are considered “homebound” if they meet these two criteria. Patients either need supportive devices such as crutches, canes, wheelchairs, and walkers; special transportation; or help from someone else in order to leave their home because of illness or injury OR have a condition that makes leaving the home medically inadvisable. There must exist a normal inability to leave home; and leaving home must require a considerable and taxing effort.

If specimen collection occurs during a home visit where other scheduled services are being provided, a specimen collection fee and travel expense cannot be billed. Payment is included in the CHHA’s per diem payment. CHHAs may bill for members who only receive lower-level services when a nurse is sent to collect the specimen (see chart above). This includes homebound patients enrolled in Managed Long-Term Care.

COVID specimen collection should not be billed to Medicaid when a home health visit is covered by Medicare (either episodic or Low Utilization Payment Adjustment (LUPA)/fee-for-service payment).

**Reporting Test Results**

Providers are required to report SARS-CoV-2 diagnostic or serology testing results, including those using SARS-CoV-2 point-of-care tests, to the Commissioner of Health through the Electronic Clinical Laboratory Reporting System (ECLRS) within 24 hours. Required reporting includes all positive, negative, and indeterminate results, as well as test type, specimen source, and several demographic details of the individual tested. Results must be reported to ECLRS through HL7 or manual entry. Contact the ECLRS Help Desk at (866) 325–7743 or via eclrs@health.ny.gov with any technical questions.

For additional testing guidance from the CDC and Wadsworth Center please see the following links:
https://coronavirus.health.ny.gov/covid-19-testing  
### Monoclonal Antibody COVID-19 Infusions

#### Payment Allowances and Effective Dates for COVID-19 Monoclonal Antibodies and their Administration

<table>
<thead>
<tr>
<th>Code</th>
<th>Labeler Name</th>
<th>Procedure Name</th>
<th>Payment Allowances</th>
<th>Effective Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Q0239</td>
<td>Eli Lilly</td>
<td>Injection, bamlanivimab, 700 mg</td>
<td>$0.00</td>
<td>11/10/2020 – 4/16/2021* EUA REVOKED</td>
</tr>
<tr>
<td>M0239</td>
<td>Eli Lilly</td>
<td>Intravenous infusion, bamlanivimab, includes infusion and post administration monitoring</td>
<td>$309.60</td>
<td>11/10/2020 – 4/16/2021* EUA REVOKED</td>
</tr>
<tr>
<td>Q0240</td>
<td>Regeneron</td>
<td>Injection, casirivimab and imdevimab, 600 mg</td>
<td>$0.00</td>
<td>7/30/2021</td>
</tr>
<tr>
<td>M0240</td>
<td>Regeneron</td>
<td>Intravenous infusion or subcutaneous injection, casirivimab and imdevimab includes infusion or injection, and post administration monitoring, subsequent repeat doses</td>
<td>$309.60</td>
<td>7/30/2021</td>
</tr>
<tr>
<td>Q0243</td>
<td>Regeneron</td>
<td>Injection, casirivimab and imdevimab, 2400 mg</td>
<td>$0.00</td>
<td>11/21/2020</td>
</tr>
<tr>
<td>Q0244</td>
<td>Regeneron</td>
<td>Injection, casirivimab and imdevimab, 1200 mg</td>
<td>$0.00</td>
<td>06/03/2021</td>
</tr>
<tr>
<td>M0243</td>
<td>Regeneron</td>
<td>Intravenous infusion, casirivimab and imdevimab, includes infusion and post administration monitoring</td>
<td>$309.60</td>
<td>11/21/2020</td>
</tr>
<tr>
<td>Q0245</td>
<td>Eli Lilly</td>
<td>Injection, bamlanivimab and etesevimab, 2100 mg</td>
<td>$0.00</td>
<td>2/09/2021</td>
</tr>
<tr>
<td>M0245</td>
<td>Eli Lilly</td>
<td>Intravenous infusion, bamlanivimab and etesevimab, includes infusion and post administration monitoring</td>
<td>$309.60</td>
<td>2/09/2021</td>
</tr>
<tr>
<td>Q0247</td>
<td>GSK</td>
<td>Injection, sotrovimab, 500 mg</td>
<td>@ Acquisition Cost per Invoice</td>
<td>05/26/2021</td>
</tr>
<tr>
<td>M0247</td>
<td>GSK</td>
<td>Intravenous infusion, sotrovimab, includes infusion and post administration monitoring</td>
<td>$309.60</td>
<td>05/26/2021</td>
</tr>
</tbody>
</table>

The above monoclonal antibody infusion codes are reimbursable when provided in a hospital outpatient department or physician’s office. Providers should bill the codes outlined in this guidance via the ordered ambulatory and/or physician fee schedule. The
Payment allowances for the above infusion administration codes include all costs for any saline, any other fluid, and/or any other drug used for the infusion and any post-infusion patient monitoring.

### Payment Allowances and Effective Dates for COVID-19 Monoclonal Antibodies Infusion Administration provided in a home setting.

<table>
<thead>
<tr>
<th>Code</th>
<th>Labeler Name</th>
<th>Procedure Name</th>
<th>Payment Allowances</th>
<th>Effective Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>M0241</td>
<td>Regeneron</td>
<td>Intravenous infusion or subcutaneous injection, casirivimab and imdevimab includes infusion or injection, and post administration monitoring in the home or residence – Subsequent repeat doses</td>
<td>$525.00</td>
<td>7/30/2021</td>
</tr>
<tr>
<td>M0244</td>
<td>Regeneron</td>
<td>Intravenous infusion or subcutaneous injection, casirivimab and imdevimab includes infusion or injection, and post administration monitoring in the home or residence</td>
<td>$525.00</td>
<td>5/6/2021</td>
</tr>
<tr>
<td>M0246</td>
<td>Eli Lilly</td>
<td>Intravenous infusion, bamlanivimab and etesevimab, includes infusion and post administration monitoring in the home or residence</td>
<td>$525.00</td>
<td>5/6/2021</td>
</tr>
<tr>
<td>M0248</td>
<td>GSK</td>
<td>Intravenous infusion, sotrovimab, includes infusion and post administration monitoring in the home or residence</td>
<td>$525.00</td>
<td>5/26/2021</td>
</tr>
</tbody>
</table>

Providers should bill the codes outlined in the table above via the physician fee schedule. The payment allowances for the above infusion administration codes include all costs for any saline, any other fluid, and/or any other drug used for the infusion and any post-infusion patient monitoring.


**Questions**
Medicaid FFS coverage and policy questions should be directed to the Office of Health Insurance Programs, Division of Program Development and Management, at (518) 473–2160 or FFSMedicaidPolicy@health.ny.gov.

MMC reimbursement, billing, and/or documentation requirement questions should be directed to the enrollee’s MMC plan.

FFS claim questions should be directed to the eMedNY Call Center at (800) 343–9000.