# Illinois Medicaid COVID-19 Fee Schedule

PLEASE NOTE: New COVID-19 related codes will be added to the HFS system as they are released by the Centers for Medicare and Medicaid Services (CMS) in accordance with the <u>December 8, 2020 provider notice</u>. Claims containing new codes which do not have a Medicare or National Government Services (NGS) rate will temporarily suspend until a code rate is assigned. Once a rate is assigned, the HFS system will be updated with that rate and any held claims released into processing.

#### **COVID-19 Vaccines and Vaccine Administration**

COVID-19 vaccine product procedure codes are included as a reference but should not be billed when obtained at zero cost to the provider. COVID-19 vaccine administration procedure codes are billable by Physicians, Advance Practice Nurses (APNs), Physician Assistants (PAs), Local Health Departments, Federally Qualified Health Centers (FQHCs), Rural Health Clinics (RHCs), Encounter Rate Clinics (ERCs), and School-Based Health Centers (SBHCs).

<u>*Please Note*</u>: FQHCs, RHCs and ERCs must submit COVID-19 vaccine administration codes fee-forservice separately from an encounter claim, even if the vaccine was administered during a face-to face encounter with a practitioner.

Procedure Code	Effective Date	Description	State Max Amount
91300	12/11/2020	Pfizer-BioNTech Covid-19 Vaccine (Aged 12 years and older) (Purple Cap)	N/A (currently government supplied at no cost to the provider)
0001A	3/15/2021 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – First Dose	42.14
0001A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – First Dose	43.60
0002A	3/15/2021 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Second Dose	42.14
0002A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Second Dose	43.60
0003A	8/12/2021 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Third Dose	42.14
0003A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Third Dose	43.60
0004A	9/22/2021 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Booster	42.14

43.6	Pfizer-BioNTech Covid-19 Vaccine (Purple Cap) Administration – Booster	1/1/2023	0004A
N/ (currently government supplie) at no cost to the provide	Moderna Covid-19 Vaccine (Aged 12 years and older) (Red Cap)	12/18/2020	91301
42.1	Moderna Covid-19 Vaccine (Red Cap) Administration – First Dose	3/15/2021 – 12/31/2022	0011A
43.6	Moderna Covid-19 Vaccine (Red Cap) Administration – First Dose	1/1/2023	0011A
42.1	Moderna Covid-19 Vaccine (Red Cap) Administration – Second Dose	3/15/2021 – 12/31/2022	0012A
43.6	Moderna Covid-19 Vaccine (Red Cap) Administration – Second Dose	1/1/2023	0012A
42.1	Moderna Covid-19 Vaccine (Red Cap) Administration – Third Dose	8/12/2021 – 12/31/2022	0013A
43.6	Moderna Covid-19 Vaccine (Red Cap) Administration – Third Dose	1/1/2023	0013A
N/ currently government supplie) at no cost to the provide	Janssen Covid-19 Vaccine (Aged 18 years and older)	2/27/2021	91303
42.1	Janssen (Johnson & Johnson) COVID- 19 Vaccine Administration – Single Dose	3/15/2021 – 12/31/2022	0031A
43.6	Janssen (Johnson & Johnson) COVID- 19 Vaccine Administration – Single Dose	1/1/2023	0031A
42.1	Janssen (Johnson & Johnson) COVID- 19 Vaccine Administration - Booster	10/20/2021 – 12/31/2022	0034A
43.6	Janssen (Johnson & Johnson) COVID- 19 Vaccine Administration - Booster	1/1/2023	0034A
N/ (currently government supplie) at no cost to the provide	Novavax Covid-19 Vaccine, Adjuvanted (Aged 18 years and older)	7/13/2022 – 8/21/2022	91304

91304	8/22/2022	Novavax Covid-19 Vaccine, Adjuvanted (Aged 12 years and older)	N/A (currently government supplied at no cost to the provider)
0041A	7/13/2022 – 12/31/2022	Novavax Covid-19 Vaccine, Adjuvanted Administration – First Dose	41.80
0041A	1/1/2023	Novavax Covid-19 Vaccine, Adjuvanted Administration – First Dose	43.60
0042A	7/13/2022 – 12/31/2022	Novavax Covid-19 Vaccine, Adjuvanted Administration – Second Dose	41.80
0042A	1/1/2023	Novavax Covid-19 Vaccine, Adjuvanted Administration – Second Dose	43.60
0044A	10/19/2022 – 12/31/2022	Novavax Covid-19 Vaccine, Adjuvanted Administration – Booster	41.80
0044A	1/1/2023	Novavax Covid-19 Vaccine, Adjuvanted Administration – Booster	43.60
91305	1/3/2022	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Aged 12 years and older) (Gray Cap)	N/A (currently government supplied at no cost to the provider)
0051A	1/3/2022 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - First dose	41.80
0051A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - First dose	43.60
0052A	1/3/2022 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Second dose	41.80
0052A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Second dose	43.60
0053A	1/3/2022 – 12/31/2022	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Third dose	41.80
0053A	1/1/2023	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Third dose	43.60

41.8	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Booster	1/3/2022 – 12/31/2022	0054A
43.6	Pfizer-BioNTech Covid-19 Vaccine Pre-Diluted (Gray Cap) Administration - Booster	1/1/2023	0054A
N/ (currently government supplie) at no cost to the provide	Moderna Covid-19 Vaccine (Aged 18 years and older) (Red Cap) (Low Dose)	10/20/2021	91306
42.1	Moderna Covid-19 Vaccine (Red Cap) (Low Dose) Administration - Booster	10/20/2021 – 12/31/2022	0064A
43.6	Moderna Covid-19 Vaccine (Red Cap) (Low Dose) Administration - Booster	1/1/2023	0064A
N/ (currently government supplie) at no cost to the provide	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Aged 5 years through 11 years) (Orange Cap)	10/29/2021	91307
42.1	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - First dose	10/29/2021 – 12/31/2022	0071A
43.6	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - First dose	1/1/2023	0071A
42.1	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Second dose	10/29/2021 – 12/31/2022	0072A
43.6	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Second dose	1/1/2023	0072A
41.8	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Third dose	01/03/2022 – 12/31/2022	0073A
43.6	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Third dose	1/1/2023	0073A
41.8	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Booster	05/17/2022 – 12/31/2022	0074A
43.6	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Orange Cap) - Administration - Booster	1/1/2023	0074A

91308	06/17/2022	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Aged 6 months through 4 years) (Maroon Cap)	N/A (currently government supplied at no cost to the provider)
0081A	06/17/2022 – 12/31/2022	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - First dose	41.80
0081A	1/1/2023	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - First dose	43.60
0082A	06/17/2022 – 12/31/2022	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - Second dose	41.80
0082A	1/1/2023	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - Second dose	43.60
0083A	06/17/2022 – 12/31/2022	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - Third dose	41.80
0083A	1/1/2023	Pfizer-BioNTech COVID-19 Pediatric Vaccine (Maroon Cap) - Administration - Third dose	43.60
91309	3/29/2022	Moderna Covid-19 Vaccine (Aged 6 years through 11 years or aged 18+) (Blue Cap with purple border) 50MCG/0.5ML	N/A (currently government supplied at no cost to the provider)
0091A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) Administration - First dose	41.80
0091A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) Administration - First dose	43.60
0092A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) - Administration - Second dose	41.80
0092A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) - Administration - Second dose	43.60
0093A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) - Administration - Third dose	41.80
0093A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with purple border) - Administration - Third dose	43.60

0094A	3/29/2022 -	Moderna Covid-19 Vaccine (Blue	41.80
	12/31/2022	Cap with purple border) 50MCG/0.5ML Administration - Booster	
0094A	1/1/2023	Moderna Covid-19 Vaccine (Blue Cap with purple border) 50MCG/0.5ML Administration - Booster	43.60
91311	6/17/2022	Moderna Covid-19 Pediatric Vaccine (Aged 6 months through 5 years) (Blue Cap with magenta border) 250MCG/0.25ML	N/A (currently government supplied at no cost to the provider)
0111A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - First dose	41.80
0111A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - First dose	43.60
0112A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - Second dose	41.80
0112A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - Second dose	43.60
0113A	6/17/2022 – 12/31/2022	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - Third dose	41.80
0113A	1/1/2023	Moderna Covid-19 Pediatric Vaccine (Blue Cap with magenta border) - Administration - Third dose	43.60
91312	8/31/2022 – 10/11/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 12 years and older) (Gray Cap)	N/A (currently government supplied at no cost to the provider)
91312	10/12/2022 – 12/7/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 5 years and older) (Gray Cap)	N/A (currently government supplied at no cost to the provider)
91312	12/8/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 6 months and older) (Gray Cap)	N/A (currently government supplied at no cost to the provider)
0124A	8/31/2022 – 12/31/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent (Gray Cap) Administration – Booster Dose	41.80
0124A	1/1/2023	Pfizer-BioNTech COVID-19 Vaccine, Bivalent (Gray Cap) Administration – Booster Dose	43.60

91313	8/31/2022 –	Moderna COVID-19 Vaccine,	N/A
	10/11/2022	Bivalent Product (Aged 18 years and older) (Dark Blue Cap with gray border)	(currently government supplied at no cost to the provider)
91313	10/12/2022 – 12/7/2022	Moderna COVID-19 Vaccine, Bivalent Product (Aged 6 years and older) (Dark Blue Cap with gray border)	N/A (currently government supplied at no cost to the provider)
91313	12/8/2022	Moderna COVID-19 Vaccine, Bivalent Product (Aged 6 months and older) (Dark Blue Cap with gray border)	N/A (currently government supplied at no cost to the provider)
0134A	8/31/2022 – 12/31/2022	Moderna COVID-19 Vaccine, Bivalent (Dark Blue Cap with gray border) Administration – Booster Dose	41.80
0134A	1/1/2023	Moderna COVID-19 Vaccine, Bivalent (Dark Blue Cap with gray border) Administration – Booster Dose	43.60
91314	10/12/2022 – 12/7/2022	Moderna COVID-19 Vaccine, Bivalent Product (Aged 6 years through 11 years) (Dark Blue Cap with gray border)	N/A (currently government supplied at no cost to the provider)
91314	12/8/2022	Moderna COVID-19 Vaccine, Bivalent Product (Aged 6 months through 11 years) (Dark Blue Cap with gray border)	N/A (currently government supplied at no cost to the provider)
0144A	10/12/2022 – 12/31/2022	Moderna COVID-19 Vaccine, Bivalent (Dark Blue Cap with gray border) Administration – Booster Dose	41.80
0144A	1/1/2023	Moderna COVID-19 Vaccine, Bivalent (Dark Blue Cap with gray border) Administration – Booster Dose	43.60
91315	10/12/2022 – 12/7/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 5 years through 11 years) (Orange Cap)	N/A (currently government supplied at no cost to the provider)
91315	12/8/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 6 months through 11 years) (Orange Cap)	N/A (currently government supplied at no cost to the provider)
0154A	10/12/2022 – 12/31/2022	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Orange Cap) Administration – Booster Dose	41.80

	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Orange Cap) Administration – Booster Dose	1/1/2023	0154A
(currently government sup at no cost to the prov	Moderna COVID-19 Vaccine, Bivalent Product (Aged 6 months through 5 years) (Dark Pink Cap and a label with a yellow box)	12/8/2022	91316
	Moderna COVID-19 Vaccine, Bivalent (Aged 6 months through 5 years) (Dark Pink Cap and label with a yellow box) Administration – Booster Dose	12/8/2022 – 12/31/2022	0164A
	Moderna COVID-19 Vaccine, Bivalent (Aged 6 months through 5 years) (Dark Pink Cap and label with a yellow box) Administration – Booster Dose	1/1/2023	0164A
(currently government sup at no cost to the prov	Pfizer-BioNTech COVID-19 Vaccine, Bivalent Product (Aged 6 months through 4 years) (Maroon Cap)	12/8/2022	91317
	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Aged 6 months through 4 years) (Maroon Cap) Administration - Third dose	12/8/2022 – 12/31/2022	0173A
	Pfizer-BioNTech Covid-19 Pediatric Vaccine (Aged 6 months through 4 years) (Maroon Cap) Administration - Third dose	1/1/2023	0173A
r	COVID-19 Vaccine Administration Inside a Patient's Home <b>Note</b> : see the HFS <u>7/2/21 provider</u> <u>notice</u> for information, though this code is no longer limited to once per DOS, per home effective 8/24/21	6/8/2021 – 12/31/2022	M0201
r	COVID-19 Vaccine Administration Inside a Patient's Home <b>Note</b> : see the HFS 7/2/21 provider notice for information, though this code is no longer limited to once per DOS, per home effective 8/24/21	1/1/2023	M0201

# Vaccine Counseling

Billable by Local Health Departments, Physicians, Advance Practice Nurses (APNs), Physician Assistants (PAs), and School-Based Health Centers (SBHCs). Intended to provide reimbursement for the additional time needed for parental/caregiver counseling and informed consent for the COVID-19 vaccination of children ages 6 months through 20. \**Note*: this code is *not billable as a telehealth service.* 

Procedure Code	Effective Date	Ages	Description	State Max Amount
99402	10/29/2021 for ages 5-20	6 months through 20	Preventive medicine counseling and/or risk factor reduction intervention(s)	30.00
	6/17/2022 for ages 6 mos–20	years	provided to an individual (separate procedure) ; approximately 30 min.	

# Virtual Healthcare/Telehealth Expansion

Billable by Local Health Departments, Physicians, Advance Practice Nurses (APNs), and Physician Assistants (PAs) – including physicians, APNs, and PAs rendering the service in a Federally Qualified Health Center (FQHC), Rural Health Clinic (RHC), Encounter Rate Clinic (ERC), or School Based Health Center (SBHC):

\**Note*: all virtual healthcare/telehealth codes must be billed with place of service 02 (or place of service 10 if applicable and date of service is on/after 7/1/2022), and modifier GT (or modifier 93 if applicable and date of service is on/after 7/1/2022).

Procedure Code	Effective Date	Description	State Max
			Amount
G0406	3/9/2020	Follow-up inpatient consultation, limited, physicians typically spend 15 minutes communicating with the patient via telehealth	39.17
G0407	3/9/2020	Follow-up inpatient consultation, intermediate, physicians typically spend 25 minutes communicating with the patient via telehealth	72.13
G0408	3/9/2020	Follow-up inpatient consultation, complex, physicians typically spend 35 minutes communicating with the patient via telehealth	103.70
G0425	3/9/2020	Telehealth consultation, emergency department or initial inpatient, typically 30 minutes communicating with the patient via telehealth	100.35
G0426	3/9/2020	Telehealth consultation, emergency department or initial inpatient, typically 50 minutes communicating with the patient via telehealth	136.14

G0427	3/9/2020	Telehealth consultation, emergency department or initial inpatient, typically 70 minutes or more communicating with the patient via telehealth	201.99
G2010	3/9/2020	Remote evaluation of recorded video and/or images submitted by an established patient (e.g., store and forward), including interpretation with follow-up with the patient within 24 business hours, not originating from a related e/m service provided within the previous 7 days nor leading to an e/m service or procedure within the next 24 hours or soonest available appointment	9.24
G2012	3/9/2020	Brief communication technology-based service, e.g. virtual check-in, by a physician or other qualified health care professional who can report evaluation and management services, provided to an established patient, not originating from a related e/m service provided within the previous 7 days nor leading to an e/m service or procedure within the next 24 hours or soonest available appointment; 5-10 minutes of medical discussion	13.19
G2061	3/9/2020 – 12/31/2020	Qualified nonphysician healthcare professional online assessment, for an established patient, for up to seven days, cumulative time during the 7 days; 5-10 minutes	12.10
G2062	3/9/2020 – 12/31/2020	Qualified nonphysician healthcare professional online assessment service, for an established patient, for up to seven days, cumulative time during the 7 days; 11-20 minutes	21.37
G2063	3/9/2020 – 12/31/2020	Qualified nonphysician qualified healthcare professional assessment service, for an established patient, for up to seven days, cumulative time during the 7 days; 21 or more minutes	33.14
G2250	1/1/2021	Remote assessment of recorded video and/or images submitted by an established patient (e.g., store and forward), including interpretation with follow-up with the patient within 24 business hours, not originating from a related service provided within the previous 7 days nor leading to a service or procedure within the next 24 hours or soonest available appointment	9.24
G2251	1/1/2021	Brief communication technology-based service, e.g. virtual check-in, by a qualified health care professional who cannot report evaluation and management services, provided to an established patient, not originating from a related service provided within the previous 7 days nor leading to a service or procedure within the next 24 hours or soonest available appointment; 5-10 minutes of clinical discussion	13.05
G2252	1/1/2021	Brief communication technology-based service, e.g. virtual check-in, by a physician or other qualified health care professional who can report evaluation and management services, provided to an established patient, not originating from a related e/m service provided within the previous 7	25.14

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ours or soonest available appointment; 11-20
of medical discussion
nonphysician health care professional online digital 11.3
nt and management, for an established patient, for
ays, cumulative time during the 7 days; 5-10 min.
nonphysician health care professional online digital 20.3
nt and management, for an established patient, for
ays, cumulative time during the 7 days; 11-20 min.
nonphysician health care professional online digital 32.4
nt and management, for an established patient, for
ays, cumulative time during the 7 days; 21+ min.
gital Evaluation and Management Service, for an 13.1
ed patient, for up to 7 days, cumulative time during
s; 5-10 minutes
gital Evaluation and Management Service, for an 27.1
ed patient, for up to 7 days, cumulative time during
s; 11-20 minutes
gital Evaluation and Management Service, for an 43.2
ed patient, for up to 7 days, cumulative time during
s; 21 or more minutes

**Please Note:** Evaluation and management services rendered by Physicians, Advance Practice Nurses, and Physician Assistants to new or existing patients using audio only telephonic equipment may be billed as a distant site telehealth service so long as the E/M service is of an amount and nature that would be sufficient to meet the key components of a face-to-face encounter. In this scenario, the claim must be submitted with place of service 02 (or 10 if applicable and the date of service is on/after 7/1/2022) and modifier GT (or 93 if applicable and the date of service is on/after 7/1/2022) appended to the applicable procedure code.

If an audio only telephonic interaction cannot meet key components of a face-to-face encounter, the provider may instead seek reimbursement for virtual check-in services using CPT code G2012. FQHCs/RHCs/ERCs will be reimbursed at the above rates (not their medical encounter rate) for virtual check-in and E-visit codes. Virtual check-in and e-visit/online portal services must be submitted fee-for-service without the T1015 encounter code.

# Virtual Healthcare/Teledentistry Expansion

**Billable by Dentists:** \*Note the below codes must be billed in addition to D0140, with place of service 02 (or 10, if applicable and date of service is on/after 7/1/2022). Do not include modifier GT or 93.

Procedure Code	Effective Date	Description	State Max Amount
D9995	3/9/2020	Teledentistry, synchronous; real-time encounter	13.19
D9996	3/9/2020	Teledentistry asynchronous; information stored and forwarded to dentist for subsequent review	9.24

#### COVID-19 Treatment

COVID-19 antibody product procedure codes are included as a reference but should not be billed when obtained at zero cost to the provider. Only Q0247 was previously practitioner-purchased until it became unauthorized effective April 5, 2022.

Antibody treatment administration codes are billable by Physicians, Advance Practice Nurses (APNs), and Physician Assistants (PAs). Federally Qualified Health Centers (FQHCs), Rural Health Clinics (RHCs), and Encounter Rate Clinics (ERCs) may bill antibody treatment administration codes as detail codes with an encounter claim.

Hospitals may bill the antibody treatment administration codes marked with a double asterisk (\*\*) using revenue code 0771. Reimbursement is based on DRG (inpatient setting) or EAPG (outpatient setting) methodology.

Procedure Code	Effective Date	Description & Labeler Name	State Max Amount
J0248	12/23/2021	Injection, REMDESIVIR, 1 mg Please reference the <u>10/21/22</u> provider notice for details	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.
Q0220	12/8/2021	Tixagev and Cilgav, 300mg	N/A (currently government supplied at no cost to the provider)
M0220	12/8/2021	Injection, Tixagevimab and Cilgavimab, for the pre-exposure prophylaxis only, for certain adults and pediatric individuals (12 years of age and older weighing at least 40kg) with no known sars-cov-2 exposure, who either have moderate to severely compromised immune systems or for whom vaccination with any available covid-19 vaccine is not recommended due to a history of severe adverse reaction to a covid- 19 vaccine(s) and/or covid-19 vaccine component(s), includes injection and post administration monitoring	150.50
Q0221	2/24/2022	Injection, Tixagevimab and Cilgavimab, for the pre-exposure prophylaxis only, for certain adults and pediatric individuals (12 years of age and older weighing at least 40kg) with no known sars-cov-2 exposure, who either have moderate to severely compromised	N/A (currently government supplied at no cost to the provider)

M0221**	12/8/2021	immune systems or for whom vaccination with any available covid-19 vaccine is not recommended due to a history of severe adverse reaction to a covid- 19 vaccine(s) and/or covid-19 vaccine component(s), 600 mg Injection, Tixagevimab and Cilgavimab, for the pre-exposure prophylaxis only, for certain adults and pediatric individuals (12 years of age and older weighing at least 40kg) with no known sars-cov-2 exposure, who either have moderate to severely compromised immune systems or for whom vaccination with any available covid-19 vaccine is not recommended due to a history of severe adverse reaction to a covid- 19 vaccine(s) and/or covid-19 vaccine component(s), includes injection and post administration monitoring in the home or residence; this includes a beneficiary's home that has been made provider-based to the hospital during the covid-19 public health emergency	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.
Q0222	2/11/2022 – 11/30/2022	Injection, Bebtelovimab, 175 mg	N/A (currently government supplied at no cost to the provider)
M0222	2/11/2022 – 11/30/2022	Intravenous injection, Bebtelovimab, includes injection and post administration monitoring	350.50
M0223**	2/11/2022 – 11/30/2022	Intravenous injection, Bebtelovimab, includes injection and post administration monitoring in the home or residence; this includes a beneficiary's home that has been made provider-based to the hospital during the covid-19 public health emergency	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.

Q0243	11/21/2020 -	Casirivimab and Imdevimab, 2400	N/A
	1/23/2022	mg (Regeneron)	(currently government supplied at no cost to the provider)
M0243**	5/6/2021 – 1/23/2022	Intravenous infusion, Casirivimab and Imdevimab, includes infusion and post administration monitoring (Regeneron)	450.00 **Hospitals billing on the 837I will be reimbursed using EAPG methodology.
M0244**	5/6/2021 – 1/23/2022	Intravenous infusion or subcutaneous injection, Casirivimab and Imdevimab includes infusion or injection, and post administration monitoring in the home or residence; this includes a beneficiary's home that has been made provider-based to the hospital during the covid-19 public health emergency	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.
Q0245	2/9/2021 – 1/23/2022	Injection, Bamlanivimab and Etesevimab, 2100 mg (Eli Lilly)	N/A (currently government supplied at no cost to the provider)
M0245	5/6/2021 – 1/23/2022	Intravenous infusion, Bamlanivimab and Etesevimab, includes infusion and post administration monitoring (Eli Lilly)	450.00
M0246**	5/6/2021 – 1/23/2022	Intravenous infusion, Bamlanivimab and Etesevimab, includes infusion and post administration monitoring in the home or residence; this includes a beneficiary's home that has been made provider-based to the hospital during the covid-19 public health emergency	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.
Q0247	10/1/2021 – 4/4/2022	Injection, Sotrovimab, 500 mg	2394.00
M0247	7/1/2021 – 4/4/2022	Intravenous infusion, Sotrovimab, includes infusion and post administration monitoring	450.00
M0248**	5/26/2021 – 4/4/2022	Intravenous infusion, Sotrovimab, includes infusion and post administration monitoring in the home or residence; this includes a beneficiary's home that has been made provider-based to the hospital during the covid-19 public health emergency	**Billable only by hospitals on the 837I. Reimbursed using EAPG methodology.

Q0249	6/24/2021	Injection, Tocilizumab, for	N/A
		hospitalized adults and pediatric	(currently government supplied
		patients (2 years of age and older)	at no cost to the provider)
		with covid-19 who are receiving	
		systemic corticosteroids and require	
		supplemental oxygen, non-invasive	
		or invasive mechanical ventilation,	
		or extracorporeal membrane	
		oxygenation (ECMO) only, 1 mg	
M0249**	6/24/2021	Intravenous infusion, Tocilizumab,	**Billable only by hospitals on
		for hospitalized adults and pediatric	the 837I. Reimbursed using DRG
		patients (2 years of age and older)	methodology.
		with covid-19 who are receiving	
		systemic corticosteroids and require	
		supplemental oxygen, non-invasive	
		or invasive mechanical ventilation,	
		or extracorporeal membrane	
		oxygenation (ECMO) only, includes	
		infusion and post administration	
		monitoring, first dose	
M0250**	6/24/2021	Intravenous infusion, Tocilizumab,	**Billable only by hospitals on
		for hospitalized adults and pediatric	the 8371. Reimbursed using DRG
		patients (2 years of age and older)	methodology.
		with covid-19 who are receiving	
		systemic corticosteroids and require	
		supplemental oxygen, non-invasive	
		or invasive mechanical ventilation,	
		or extracorporeal membrane	
		oxygenation (ECMO) only, includes	
		infusion and post administration	
		monitoring, second dose	

## **Laboratory Services**

Billable by Local Health Departments, Physicians, Advance Practice Nurses (APNs), Physician Assistants (PAs), and Independent Laboratories. Federally Qualified Health Centers (FQHCs), Rural Health Clinics (RHCs), and Encounter Rate Clinics (ERCs) may bill the following laboratory services as detail codes with an encounter claim when the laboratory service is performed on-site. Please note that providers normally subject to a SMART Act rate reduction (e.g. Independent Labs) must include modifier CR to exempt the service from a rate reduction.

Hospitals must bill on an institutional invoice and will be reimbursed via the EAPG methodology.

Procedure	Effective	Description	State
Code	Date		Max
			Amount

0202U	5/20/2020	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	250.07
0223U	6/25/2020	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	416.78
0224U	6/25/2020	Antibody, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]), includes titer(s), when performed	42.13
0225U	8/10/2020	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	416.78
0226U	8/10/2020	Surrogate viral neutralization test (sVNT), severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19]), ELISA, plasma, serum	42.28
0240U	10/6/2020	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	142.63
0241U	10/6/2020	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	142.63
86318	4/10/2020	Immunoassay for infectious agent antibody(ies), qualitative or semiqualitative, single step method (e.g. reagent strip)	16.90
86328	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])	45.23
86408	8/10/2020	Neutralizing antibody, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID- 19]); screen	42.13

86409	8/10/2020	Neutralizing antibody, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID- 19]); titer	105.33
86413	9/8/2020	Severe acute respiratory syndrome coronavirus 2 (SARS-CoV- 2) (Coronavirus disease [COVID-19]) antibody, quantitative	51.43
86769	4/10/2020	Antibody; severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19])	42.13
87426	6/25/2020	Infectious agent antigen detection by immunoassay technique, (eg, enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative, multiple-step method; severe acute respiratory syndrome coronavirus (eg, SARS-CoV, SARS-CoV-2 [COVID-19])	35.33
87428	11/10/2020	Infectious agent antigen detection by immunoassay technique, (eg, enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], fluorescence immunoassay [FIA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative; severe acute respiratory syndrome coronavirus (eg, SARS-CoV, SARS-CoV-2 [COVID-19]) and influenza virus types A and B SD: SARSCOV & INF VIR A&B AG IA	63.59
87635	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	51.31
87636	10/6/2020	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	142.63
87637	10/6/2020	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	142.63
87811	10/6/2020	Infectious agent antigen detection by immunoassay with direct optical (ie, visual) observation; severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID-19])	41.38
U0001	2/4/2020	CDC 2019-Novel Coronavirus real-time RT-PCR diagnostic panel	35.91
U0002	2/4/2020	Coronavirus (COVID-19) SARS-COV-2/2019-NCOV, Non-CDC Lab Test	51.31
U0003	4/14/2020 through 2/28/2021	Infectious Agent Detection by Nucleic Acid (DNA or RNA); SARS-COV-2, COVID-19, Amplified Probe Technique, High Throughput Technologies	100.00

U0003	3/1/2021	Infectious Agent Detection by Nucleic Acid (DNA or RNA);	75.00
		SARS-COV-2, COVID-19, Amplified Probe Technique, High	
		Throughput Technologies	
U0004	4/14/2020	2019-NCOV Coronavirus, SARS-CoV-2/2019-nCoV (COVID-	100.00
	through	19), Any Technique, Multiple Subtypes, Non-CDC, High	
	2/28/2021	Throughput Technologies	
U0004	3/1/2021	2019-NCOV Coronavirus, SARS-CoV-2/2019-nCoV (COVID-	75.00
		19), specimen collection. ( <sup>+</sup> add-on to U0003 or U0004 Any	
		Technique, Multiple Subtypes, Non-CDC, High Throughput	
		Technologies	
+U0005	3/1/2021	Infectious agent detection by nucleic acid (DNA or RNA);	25.00
		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	
		and time of; List separately in addition to either HCPCS code	
		U0003 or U0004)	
		<u>NOTE:</u> certain conditions must be met to bill this code ;	
		refer to the 02/26/2021 provider notice for billing	
		guidelines	

## **COVID-19 Diagnostic Testing Specimen Collection**

\*Billable by Local Health Departments, Physicians, Advance Practice Nurses (APNs), Physician Assistants (PAs), and Federally Qualified Health Centers (FQHCs) with drive-thru testing sites. FQHCs may bill fee-for-service when there is not a billable medical encounter. Please note providers normally subject to a SMART Act rate reduction (e.g. Independent Labs) must include modifier CR to exempt the service from a rate reduction.

\*\*Billable by Local Health Departments, Physicians, Advance Practice Nurses (APNs), Physician Assistants (PAs), and Independent Labs.

Procedure	Effective	Description	State
Code	Date		Max
			Amount
99000*	3/18/2020	Handling of Specimen for Transfer from Office to a Lab	23.46
G2023**	3/1/2020	Specimen Collection, SARS-CoV-2, COVID-19, any specimen source	23.46
G2024***	3/1/2020	Specimen collection for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (Coronavirus disease [COVID- 19]), from an individual in a skilled nursing facility or by a laboratory on behalf of a home health agency, any specimen source	25.46

## <u>COVID-19 Testing, Testing-Related and Vaccination Coverage for the Uninsured</u> <u>Population</u>

The following procedure codes are covered for the uninsured population for the purposes of COVID-19 testing, testing-related services, and vaccination for dates of service beginning March 18, 2020. Testing-related services include those directly related to the administration of an in vitro diagnostic product described in section 1905(a)(3)(B) of the Social Security Act, or to the evaluation of a patient for purposes of determining the need for such product.

- HCPCS codes: G2010, G2012, G2023, G2024, G2061, G2062, G2063, G2250, G2251, G2252, T1015
- CPT codes: All COVID-19 laboratory testing and vaccine administration codes\*, 71045, 71046, 71047, 71048, 99000, 99201 (note this code became obsolete 1/1/2021), 99202, 99203, 99204, 99205, 99211, 99212, 99213, 99214, 99215, 99421, 99422, 99423
- All COVID-19 laboratory testing and vaccine administration codes

## **COVID-19 Treatment Coverage for the Uninsured Population**

In accordance with the <u>December 22, 2022 provider notice</u>, practitioners may now submit professional claims for monoclonal antibody treatment for a specific subset of the uninsured population via the provider's usual claim submittal process **outside the HFS Uninsured Portal**. In order to identify the uninsured individuals eligible for monoclonal antibody treatment:

- 1) Check the MEDI system to verify if the person has an existing Recipient Identification Number (RIN) assigned with 'COVID 19 Testing Only' eligibility for the date of service.
- 2) If a RIN is found, the "Special Information" under the "COVID-19 Testing" eligibility in MEDI will show "Title XIX". Please note, if the "Special Information" shows only 'State-Funded' information, then COVID-19 treatment is not a covered service for that individual.

Rates for the COVID-19 testing, testing-related, vaccination and treatment services above may be found on the COVID-19 Fee Schedule preceding this uninsured coverage information, on the <u>Practitioner Fee Schedule</u> or, in the case of T1015 will be at the FQHC/RHC/ERC provider-specific medical encounter rate. Coverage effective dates are specific to each procedure code's effective date as indicated.

Providers normally subject to a SMART Act rate reduction must include modifier CR to exempt the COVID-related service from the rate reduction (e.g. independent labs billing for testing).

PLEASE NOTE: All claims for the uninsured population must contain a diagnosis code indicating the patient encounter was for the purposes of COVID testing, COVID vaccine administration, or COVID treatment. Paid claims with no COVID or COVID-related diagnosis code are subject to post-payment review and recoupment.