

NEBRASKA MEDICAID AD HOC FEE SCHEDULE

**This fee schedule is subject to change, as further Center of Medicare & Medicaid's (CMS) guidance occurs.
The updates in this version are reflected in red.**

COVID-19 CODES

| CODE | MOD | DESCRIPTION | PA | COMMENTS | COPAY | MEDICAID ALLOWABLE EFFECTIVE FOR CLAIMS WITH DOS THROUGH TO 3/14/2021 | MEDICAID ALLOWABLE EFFECTIVE FOR CLAIMS WITH DOS ON OR AFTER 3/15/2021 |
|-------|-----|--|----|---|-------|---|--|
| 0031A | | JANSEN SINGLE DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRAT | | RATE CHANGE \$36.94 Effective 1/1/2022 | | \$28.39 | \$37.25 |
| 0034A | | JANSEN COVID-19 VACCINE ADMINISTRATION -BOOSTER | | RATE CHANGE \$36.94 Effective 1/1/2022 | | | \$37.25 Effective 11/2/2021 |
| 0041A | | Novavax Covid-19 Vaccine, Adjuvanted Administration – First Dose ‘Other’ ADM SARSCOV2 5MCG/0.5ML 1ST | | | | | \$36.94 Effective 7/13/2022 |
| 0042A | | Novavax Covid-19 Vaccine, Adjuvanted Administration – Second Dose ‘Other’ ADM SARSCOV2 5MCG/0.5ML 2ND | | | | | \$36.94 Effective 7/13/2022 |
| 0051A | | Pfizer-BioNTech Covid-19 Vaccine (Ready to Use) Administration - First dose | | | | | \$36.94 Effective 12/27/2021 |
| 0052A | | Pfizer-BioNTech Covid-19 Vaccine (Ready to Use) Administration - Second dose | | | | | \$36.94 Effective 12/27/2021 |
| 0053A | | Pfizer-BioNTech Covid-19 Vaccine (Ready to Use) Administration - Third dose | | | | | \$36.94 Effective 12/27/2021 |
| 0054A | | Pfizer-BioNTech Covid-19 Vaccine (Ready to Use) Administration - Booster | | | | | \$36.94 Effective 12/27/2021 |
| 0071A | | Pfizer-BioNTech Covid-19 Pediatric Vaccine - Administration - First dose | | RATE CHANGE \$36.94 Effective 1/1/2022 | | | \$37.25 Effective 11/2/2021 |
| 0072A | | Pfizer-BioNTech Covid-19 Pediatric Vaccine - Administration - Second dose | | RATE CHANGE \$36.94 Effective 1/1/2022 | | | \$37.25 Effective 11/2/2021 |
| 0081A | | Pfizer-6 months through 4 years: Severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 3 mcg/0.2 mL dosage, diluent reconstituted, trisucrose formulation, for intramuscular use - 1st Dose | | | | | \$36.94 Effective 6/17/2022 |
| 0082A | | Pfizer-6 months through 4 years: Severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 3 mcg/0.2 mL dosage, diluent reconstituted, trisucrose formulation, for intramuscular use - 2nd Dose | | | | | \$36.94 Effective 6/17/2022 |
| 0083A | | Pfizer-6 months through 4 years: Severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 3 mcg/0.2 mL dosage, diluent reconstituted, trisucrose formulation, for intramuscular use - 3rd Dose | | | | | \$36.94 Effective 6/17/2022 |
| 0091A | | MODERNA-Immunization administration by intramuscular injection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 50 mcg/0.5 mL dosage; first dose, when administered to individuals 6 through 11 years: ADM SARSCOV2 50 MCG/.5 ML1ST | | | | | \$36.94 Effective 6/17/2022 |
| 0092A | | MODERNA- Immunization administration by intramuscular injection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 50 mcg/0.5 mL dosage; second dose, when administered to individuals 6 through 11 years: ADM SARSCOV2 50 MCG/.5 ML2ND | | | | | \$36.94 Effective 6/17/2022 |
| 0093A | | MODERNA-Immunization administration by intramuscular injection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 50 mcg/0.5 mL dosage; third dose, when administered to individuals 6 through 11 years: : ADM SARSCOV2 50 MCG/.5 ML3RD | | | | | \$36.94 Effective 6/17/2022 |
| 0111A | | Moderna-6 months through 5 years: Severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP spike protein preservative free, 25 mcg/0.25 mL dosage, for intramuscular use - 1st Dose, ADM SARSCOV2 25MCG/0.25ML1ST | | | | | \$36.94 Effective 6/17/2022 |

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| 0112A | | Moderna-6 months through 5 years: Severe acute respiratory syndrome coronavirus 2 (SARSCoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP spike protein preservative free, 25 mcg/0.25 mL dosage, for intramuscular use - 2nd Dose, ADM SARSCOV2 25MCG/0.25ML2ND | | | | | \$36.94 Effective 6/17/2022 |
| 0113A | | Moderna: Immunization administration by intramuscular injection of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (coronavirus disease [COVID-19]) vaccine, mRNA-LNP, spike protein, preservative free, 25 mcg/0.25 mL dosage; third dose, ADM SARSCOV2 25MCG/0.25ML3RD | | | | | \$36.94 Effective 6/17/2022 |
| 0124A | | PFIZER-BIONTECH COVID-19 VACCINE, BIVALENT (GRAYCAP) ADMINISTRATION BOOSTER DOSE-ADM SARSCV2 BVL 30MCG/.3ML B | | | | CODE NOT ACTIVE DURING THIS TIME | \$36.94 EFFECTIVE 08/31/2022 |
| 0134A | | MODERNA COVID-19 VACCINE, BIVALENT (AGED 18 YEARS AND OLDER) (DARK BLUE CAP WITH GRAY BORDER) ADMINISTRATION BOOSTER DOSE ADM SARSCV2 BVL 50MCG/.5ML B | | | | CODE NOT ACTIVE DURING THIS TIME | \$36.94 EFFECTIVE 08/31/2022 |
| 0073A | | IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OR SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE) | | | | | \$36.94 Effective 1/4/2022 |
| 0074A | | PFIZER-BIONTECH COVID-19 PEDIATRIC VACCINE (ORANGE CAP) ADMINISTRATION BOOSTER-ADM SARSCV2 10MCG TRS-SUCRB | | | | | \$36.94 Effective 5/17/2022 |
| 0094A | | MODERNA-ADM SARSCOV2 50MCG/0.5 MLBST | | | | | \$36.94 Effective 3/29/2022 |
| 0004A | | Pfizer-BioNTech Covid-19 Vaccine Administration – Booster | | | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$37.25 Effective 9/22/2021 |
| 0064A | | Moderna Covid-19 Vaccine (Low Dose) Administration - Booster | | | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$37.25 Effective 10/20/2021 |
| 0003A | | PFIZER BIONTECH COVID-19 ADM SARSCOV2 30MCG/0.3ML 3RD DOSE | | | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$37.25 Effective 8/12/2021 |
| 0013A | | MODERNA COVID-19 ADM SARSCOV2 100MCG/0.5ML 3RD DOSE: EFFECTIVE FOR AGE 12 AND OLDER 6-17-2022 | | | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$37.25 Effective 8/12/2021 |
| M0201 | | COVID-19 VACCINEADMINISTRATION INSIDE A PATIENT'S HOME | | | | RATE CHANGE \$33.05 Effective 1/1/2022 | \$32.50 Effective 8/6/2021 |
| M0220 | | INJECTION, TIXAGENIVIMAB AND CILGAVIMAB, FOR THE PRE-EXPOSURE PROPHYLAXIS ONLY, FOR CERTAIN ADULTS AND PEDIATRIC INDIVIDUALS; INJECTION AND ADMINISTRATION MONITORING. | | | | | \$135.31 Effective 12/8/2021 |
| M0221 | | INJECTION, TIXAGENIVIMAB AND CILGAVIMAB, FOR THE PRE-EXPOSURE PROPHYLAXIS ONLY, FOR CERTAIN ADULTS AND PEDIATRIC INDIVIDUALS; INJECTION AND ADMINISTRATION MONITORING IN THE HOME OR RESIDENCE | | | | | \$225.36 Effective 12/8/2021 |
| M0222 | | INTRAVENOUS INJECTION, BEBTELOVIMAB. INCLUDES INJECTION AND POST ADMINISTRATION MONITORING | | | | | \$350.50 Effective 2/11/2022 |
| M0223 | | 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 | | | | | \$550.50 Effective 2/11/2022 |
| Q0221 | | 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) 600mg | | | | | Not covered while federally supplied Effective 2/24/2022 |
| Q0222 | | INJECTION, BEBTELOVIAMAB, 175MG | | | | | Not covered while federally supplied Effective 2/11/2022 |

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| M0240 | | INTRAVENOUS INFUSION OR SUBCUTANEOUS INJECTION, CASIRIVIMAB AND IMDEVIMAB INCLUDES INFUSION OR INJECTION, AND POST ADMINISTRATION MONITORING, SUBSEQUENT REPEAT DOSES. | | RATE CHANGE \$404.92 Effective 12/8/2021 | | \$404.78 Effective 7/30/2021 |
| M0241 | | INTRAVENOUS INFUSION OR SUBCUTANEOUS INJECTION, CASIRIVIMAB AND IMDEVIMAB INCLUDES INFUSION OR INJECTION, AND POST ADMINISTRATION MONITORING IN THE HOME OR RESIDENCE, SUBSEQUENT REPEAT DOSES. | | RATE CHANGE \$674.52 Effective 12/8/2021 | | \$674.94 Effective 7/30/2021 |
| M0244 | | INTRAVENOUS INFUSION, CASIRIVIMAB AND IMDEVIMAB INCLUDES INFUSION AND POST ADMINISTRATION MONITORING | | RATE CHANGE \$674.52 Effective 12/8/2021 | | \$674.94 Effective 5/6/2021 |
| M0246 | | INTRAVENOUS INFUSION, BAMLANIVIMAB AND ETESEVIMAB, INCLUDES INFUSION AND POST ADMINISTRATION MONITORING IN THE HOME OR RESIDENCE DURING COVID-19 PHE | | RATE CHANGE \$674.52 Effective 12/8/2021 | | \$674.94 Effective 5/6/2021 |
| M0247 | | INTRAVENOUS INFUSION, SOTROVIMAB, INCLUDES INFUSION AND POST ADMINISTRATION MONITORING | | RATE CHANGE \$404.92 Effective 12/8/2021 | | \$404.78 Effective 5/26/2021 |
| M0248 | | INTRAVENOUS INFUSION, SOTROVIMAB, INCLUDES INFUSION AND POST ADMINISTRATION MONITORING IN THE HOME OR RESIDENCE | | RATE CHANGE \$674.52 Effective 12/8/2021 | | \$674.94 Effective 5/26/2021 |
| M0249 | | INTRAVENOUS INFUSION, TOCILIZUMAB, FOR HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS.. 1ST DOSE | | RATE CHANGE \$404.92 Effective 12/8/2021 | | \$404.78 Effective 6/24/2021 |
| M0250 | | INTRAVENOUS INFUSION, TOCILIZUMAB, FOR HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS.. 2ND DOSE | | RATE CHANGE \$404.92 Effective 12/8/2021 | | \$404.78 Effective 6/24/2021 |
| Q0247 | | INJECTION, SOTROVIMAB, 500mg | | | | \$2,394.00 Effective 5/26/2021 |
| Q0249 | | INJECTION, TOCILIZUMAB, FOR HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS.. WITH COVID-19...ECMO ONLY 1 MG | | | | \$6.57 Effective 6/24/2021 |
| D0606 | | MOLECULAR TESTING FOR A PUBLIC HEALTH RELATED PATHOGEN, INCLUDING CORONA VIRUS | | RATE CHANGE \$56.44 EFFECTIVE 7/1/2022 | | \$51.31 |
| D0171 | | RE-EVALUATION, POST-OPERATIVE OFFICE VISIT | | | | \$17.95 Effective 7/1/2022 |
| D1701 | | PFIZER 1ST DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY INFECTION | | RATE CHANGE \$40.98 EFFECTIVE 7/1/2022 | | \$37.25 |
| D1702 | | PFIZER 2ND DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY INFECTION | | RATE CHANGE \$40.98 EFFECTIVE 7/1/2022 | | \$37.25 |
| D1703 | | MODERNA 1ST DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY INFECTION | | RATE CHANGE \$40.98 EFFECTIVE 7/1/2022 | | \$37.25 |
| D1704 | | MODERNA 2ND DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY INFECTION | | RATE CHANGE \$40.98 EFFECTIVE 7/1/2022 | | \$37.25 |
| D1707 | | JANSSEN SINGLE DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY INFECTION | | RATE CHANGE \$40.98 EFFECTIVE 7/1/2022 | | \$37.25 |
| D1708 | | Pfizer-biontech covid-19 vaccine administration - third dose | | | | \$40.98 Effective 7/1/2022 |
| D1709 | | Pfizer-biontech covid-19 vaccine administration - booster dose | | | | \$40.98 Effective 7/1/2022 |
| D1710 | | Moderna covid-19 vaccine administration - third dose | | | | \$40.98 Effective 7/1/2022 |
| D1711 | | Moderna covid-19 vaccine administration - booster dose | | | | \$40.98 Effective 7/1/2022 |
| D1712 | | Janssen covid-19 vaccine administration - booster dose | | | | \$40.98 Effective 7/1/2022 |
| D1713 | | Pfizer-biontech covid-19 vaccine administration tris-sucrose pediatric - first dose | | | | \$40.98 Effective 7/1/2022 |
| D1714 | | Pfizer-biontech covid-19 vaccine administration tris-sucrose pediatric - second dose | | | | \$40.98 Effective 7/1/2022 |
| D9999 | | Unspecified Adjunctive procedure | | | | \$15.92 Effective 7/1/2022 |
| 87426 | QW | INFECTIOUS AGENT ANTIGEN DETECTION BY IMMUNOASSAY TECHNIQUE, (EG, ENZYME IMMUNOASSAY, ENZYME-LINKED IMMUNOSORBENT ASSAY, | | COVID EMERGENCY PERIOD | \$35.33 | |
| 87428 | QW | INFECTIOUS AGENT ANTIGEN DETECTION BY IMMUNOASSAY TECHNIQUE, (EG, ENZYME IMMUNOASSAY ENZYME-LINKED, IMMUNOSORBENT ASSAY, FLUORESC | | COVID EMERGENCY PERIOD | \$63.59 | \$30.94 EFFECTIVE 1/1/2022 |
| 87635 | QW | AMPLIFIED DNA OR RNA PROBE DETECTION OF SARS CORONAVIRUS | | COVID EMERGENCY PERIOD | \$51.31 | |

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| M0243 | | INTRAVENOUS INFUSION, CASIRIVIMAB AND IMDEVIMAB INCLUDES INFUSION AND POST ADMINISTRATION MONITORING | | RATE CHANGE \$404.92 Effective 12/08/2021 | \$309.60 | \$404.78 Effective 5/6/2021 |
| M0245 | | INTRAVENOUS INFUSION, BAMLANIVIMAB AND ETESEVIMAB, INCLUDES INFUSION AND POST ADMINISTRATION MONITORING | | RATE CHANGE \$404.92 Effective 12/08/2021 | \$309.60 | \$404.78 Effective 5/6/2021 |
| C9803 | | HOSPITAL OUTPATIENT CLINIC VISIT SPECIMEN COLLECTION FOR (SARS-COV-2) (COVID-19), ANY SPECIMEN SOURCE | | COVID EMERGENCY PERIOD | \$23.00 | |
| G2023 | | SPECIMEN COLLECTION FOR COVID-19, ANY SPECIMEN SOURCE | | COVID EMERGENCY PERIOD | \$23.46 | |
| G2024 | | SPECIMEN COLLECTION FOR COVID-19, FROM AN INDIVIDUAL IN A SNF OR LABORATORY ON BEHALF OF A HOME HEALTH AGENCY, ANY SPECIMEN SOURCE | | COVID EMERGENCY PERIOD | \$25.46 | |
| U0001 | | CDC 2019 NOVEL CORONAVIRUS (2019-NCOV) REAL-TIME RT-PCR DIAGNOSTIC PANEL | | COVID EMERGENCY PERIOD | \$35.92 | |
| U0002 | | NON-CDC SARS-COV-2/2019-NCOV (COVID-19) | | COVID EMERGENCY PERIOD | \$51.31 | |
| U0002 | QW | NON-CDC SARS -COV-2/2019 -NCOV | | COVID EMERGENCY PERIOD | \$51.31 | |
| U0003 | | INFECTIOUS AGENT DETECTION BY DNA OR RNA;COVID-19,AMPLIFIED PROBE TECHNIQUE,HIGH THROUGHPUT TECHNOLOGIES AS DESCRIBED BY CMS-2020-01-R | | COVID EMERGENCY PERIOD | \$100.00 | \$75.00 |
| U0004 | | COVID-19,ANY TECHNIQUE,MULTIPLE TYPES OR SUBTYPES,NON-CDC, MAKING USE OF HIGH THROUGHPUT TECHNOLOGIES AS DESCRIBED BY CMS-2020-01-R | | COVID EMERGENCY PERIOD | \$100.00 | \$75.00 |
| U0005 | | INFECTIOUS AGENT DETECTION BY NUCLEIC ACID (DNA OR RNA); SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE YCO | | COVID EMERGENCY PERIOD | \$25.00 | |
| 0001A | | PFIZER 1ST DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$16.94 | \$37.25 |
| 0002A | | PFIZER 2ND DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$28.39 | \$37.25 |
| 0011A | | BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY: EFFECTIVE FOR AGE 12 AND OLDER 6/17/2022 | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$16.94 | \$37.25 |
| 0012A | | MODERNA 2ND DOSE: IMMUNIZATION ADMINISTRATION BY INTRAMUSCULAR INJECTION OF SEVERE ACUTE RESPIRATORY: EFFECTIVE FOR AGE 12 AND OLDER 6/17/2022 | | RATE CHANGE \$36.94 Effective 1/1/2022 | \$28.39 | \$37.25 |
| 0202U | | INFECTIOUS DISEASE(BACTERIAL OR VIRAL RESPIRATORY TRACT INFECTION), PATHOGEN-SPECIFIC NUCLEIC ACID (DNA OR RNA), 22 TARGETS INCLUDING SEVERE A | | COVID EMERGENCY PERIOD | \$298.60 | \$416.78 EFFECTIVE 1/1/2021 |
| 0223U | | INFECTIOUS DISEASE(BACTERIAL OR VIRAL RESPIRATORY TRACT INFECTION), PATHOGEN-SPECIFIC NUCLEIC ACID (DNA OR RNA), 22 TARGETS INCLUDING SEVERE A | | COVID EMERGENCY PERIOD | \$298.60 | \$416.78 EFFECTIVE 1/1/2021 |
| 0224U | | ANTIBODY, SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2(SARS-COV-2) (CORONAVIRUS DISEASE YCOVID-19"), INCLUDES TITER(S), WHEN PERFORMED (DO | | COVID EMERGENCY PERIOD | \$42.13 | |
| 0225U | | INFECTIOUS DISEASE (BACTERIAL OR VIRAL RESPIRATORY TRACT INFECTION) PATHOGEN-SPECIFIC DNA AND RNA, 21 TARGETS, INCLUDING SEVERE ACUTE RESPIRAT | | COVID EMERGENCY PERIOD | \$298.60 | \$416.78 EFFECTIVE 1/1/2021 |
| 0226U | | SURROGATE VIRAL NEUTRALIZATION TEST (SVNT), SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE YCOVID-19"), ELI | | COVID EMERGENCY PERIOD | \$42.28 | |
| 0240U | | INFECTIOUS DISEASE (VIRAL RESPIRATORY TRACT INFECTION),PATHOGEN-SPECIFIC RNA, 3 TARGETS (SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 SARS) | | COVID EMERGENCY PERIOD | \$142.63 | |
| 0240U | QW | INFECTIOUS DISEASE (VIRAL RESPIRATORY TRACT INFECTION),PATHOGEN-SPECIFIC RNA, 3 TARGETS (SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 SARS) | | | | \$142.63 EFFECTIVE 1/1/2022 |
| 0241U | | INFECTIOUS DISEASE (VIRAL RESPIRATORY TRACT INFECTION),PATHOGEN-SPECIFIC RNA, 4 TARGETS (SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 SARS) | | COVID EMERGENCY PERIOD | \$142.63 | |
| 0241U | QW | INFECTIOUS DISEASE (VIRAL RESPIRATORY TRACT INFECTION),PATHOGEN-SPECIFIC RNA, 4 TARGETS (SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 SARS) | | | | \$142.63 EFFECTIVE 1/1/2022 |
| 86328 | | IMMUNOASSAY FOR INFECTIOUS AGENT ANTIBODY(IES), QUALITATIVE OR SEMIQUANTITATIVE, SINGLE STEP METHOD (EG, REAGENT STRIP); (SARS-COV-2)(COVID-19) | | COVID EMERGENCY PERIOD | \$45.23 | \$45.28 EFFECTIVE 1/1/2022 |
| 86328 | QW | TEST FOR DETECTION OF SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 COVID-19 | | | | \$45.28 EFFECTIVE 1/1/2022 |
| 86408 | | SCREENING TEST FOR DETECTION OF SEVERE ACUTE RESPIRATORY SUNDROME CORONAVIRUS 2 | | COVID EMERGENCY PERIOD | \$42.13 | |

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| 86409 | | MEASUREMENT OF NEUTRALIZING ANTIBODY TO SEVERE ACUTE RESPIRATORY SYNDROME CORONA | | COVID EMERGENCY PERIOD | | \$79.61 | |
| 86769 | | ANTIBODY; SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2(SARS-COV-2) (COVID-19) | | COVID EMERGENCY PERIOD | | \$42.13 | |
| 87301 | | INFECTIOUS AGENT ANTIGEN DETECTION BY ENZYME IMMUNOASSAY TECHNIQUE, QUALITATIVE OR SEMIQUANTITATIVE, MULTIPLE STEP METHOD; ADENO VIRUS | | COVID EMERGENCY PERIOD | | \$11.98 | |
| 87426 | | INFECTIOUS AGENT ANTIGEN DETECTION BY IMMUNOASSAY TECHNIQUE,(EG, ENZYME IMMUNOASSAY YÉIA ", ENZYME-LINKED IMMUNOSORBENT ASSAY YÉLISA ", IMMUNOCH | | COVID EMERGENCY PERIOD | | \$35.33 | |
| 87428 | | INFECTIOUS AGENT ANTIGEN DETECTION BY IMMUNOASSAY TECHNIQUE,(EG, ENZYME IMMUNOASSAY YÉIA ", ENZYME-LINKED IMMUNOSORBENT ASSAY YÉLISA ", FLUORESC | | COVID EMERGENCY PERIOD | | \$63.59 | \$30.94 EFFECTIVE 1/1/2022 |
| 87635 | | DETECTION OF SARS-COV-2 AND ANY PAN-CORONAVIRUS TYPES OR SUBTYPES;INFECTIOUS AGENT DETECTION BY NUCLEIC ACID(DNA OR RNA; SEVERE ACUTE RESPIRATOR | | COVID EMERGENCY PERIOD | | \$51.31 | |
| 87636 | | INFECTIOUS AGENT DETECTION BY NUCLEIC ACID (DNA OR RNA); SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE YCO | | COVID EMERGENCY PERIOD | | \$142.63 | |
| 87636 | QW | DETECTION TEST BY MULTIPLEX AMPLIFIED PROBE TECHNIQUE FOR SEVERE ACUTE RESPIRATORY VIRUS | | | | | \$142.63 EFFECTIVE 1/1/2022 |
| 87637 | | INFECTIOUS AGENT DETECTION BY NUCLEIC ACID (DNA OR RNA); SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE YCO | | COVID EMERGENCY PERIOD | | \$142.63 | |
| 87637 | QW | DETECTION TEST BY MULTIPLEX AMPLIFIED PROBE TECHNIQUE FOR SEVERE ACUTE RESPIRATORY VIRUS | | | | | \$142.63 EFFECTIVE 1/1/2022 |
| 87811 | | INFECTIOUS AGENT DETECTION BY NUCLEIC ACID (DNA OR RNA); SEVERE ACUTE RESPIRATORY SYNDROME CORONAVIRUS 2 (SARS-COV-2) (CORONAVIRUS DISEASE YCO | | COVID EMERGENCY PERIOD | | \$41.38 | |