

List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)

A new technology for COVID-19 detection has become available that is much simpler and faster to perform than currently-recommended nucleic acid amplification tests (NAAT), like PCR. This method relies on direct detection of SARS-CoV-2 viral proteins in nasal swabs and other respiratory secretions using a lateral flow immunoassay (also called an RDT) that gives results in < 30 minutes. Though these antigen detection RDTs (Ag-RDTs) are substantially less sensitive than NAAT, they offer the possibility of rapid, inexpensive and early detection of the most infectious COVID-19 cases in appropriate settings. For more detailed technical advise please consult the WHO Interim guidance available at: <https://www.who.int/publications/i/item/antigen-detection-in-the-diagnosis-of-sars-cov-2infection-using-rapid-immunoassays>

SARS-CoV-2 Antigen Rapid Diagnostic Tests
(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|--|---|-------------------------|-------------------------------------|---|---|---------------------------------|---------------------------------|---------------|--|---------------------------------|
| 195-000 | BinaxNOW COVID-19 Ag Card | 40T/kit | n/a | Abbott Diagnostics Scarborough, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 22 months | see IFU | see IFU | Visual read | US FDA EUA |
| 41FK10 41FK20 | Panbio COVID-19 Ag Rapid Test Device (NASOPHARYNGEAL) | 25T/kit | n/a | Abbott Rapid Diagnostics Jena GmbH | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | see IFU | Visual read | TGA WHO EUL |
| 41FK11 41FK21 | Panbio COVID-19 Ag Rapid Test Device (NASAL) | 25T/kit | n/a | Abbott Rapid Diagnostics Jena GmbH | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | nasal | For consumables and details of componants refer to IFU | Health Canada TGA WHO EUL |
| Lo31-125Q5 | Flowflex SARS-CoV-2 Antigen Rapid Test | | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | nasal | Visual read | US FDA EUA |
| Lo31-11815 Lo31-125V5 Lo31-129Z5 | Flowflex SARS-CoV-2 Antigen Rapid Test | | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | SG | see IFU | Nasopharyngal | | Health Canada |
| Lo31-129R5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|-------------------------------------|---|---|---------------------------------|---------------------------------|----------------|----------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| L031-129T5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |
| L031-129U5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 5 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |
| L031-129V5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasopharyngeal | | WHO EUL |
| L031-129W5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasopharyngeal | | WHO EUL |
| L031-129Y5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 5 T/kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |
| L031-129K5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/Kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |
| L031-129M5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 25 T/Kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasopharyngeal | | WHO EUL |
| L031-129L5 | Flowflex SARS-CoV-2 Antigen Rapid Test | 5 T/Kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasal | | WHO EUL |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|----------------|--|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| Lo31-129N6 | Flowflex SARS-CoV-2 Antigen Rapid Test | 5 T/Kit | n/a | Acon Biotech (Hangzhou) Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasopharyngeal | | WHO EUL |
| | Arsonic COVID-19 Ag | | n/a | Alfresa Pharma Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | PMDA |
| PN-0003KT40 | NIDS COVID-19 Antigen Rapid Test Kit | 40T/kit | n/a | ANP Technologies, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| 1066-40 | OSOM COVID-19 Antigen Rapid Test | | n/a | ANP Technologies, Inc. (distributed by SEKISUI Diagnostics LLC) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of components refer to IFU | US FDA EUA |
| | ARISTA™ COVID-19 Antigen Rapid Test | | n/a | Arista Biotech Pte Ltd (Singapore) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| A03-50-422 | Artron COVID-19 Antigen Test | 25T/kit | n/a | Artron Laboratories Inc. (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 18 | see IFU | see IFU | | Health Canada WHO EUL |
| | COVID-19 Antigen Rapid Test Device | 25T/kit | n/a | Assure Tech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | Health Canada TGA |
| | Ecotest COVID-19 Antigen Saliva Test Kit | | n/a | Assure Tech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|---|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Ecotest COVID-19 Antigen Nasal Test Kit | | n/a | Assure Tech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| | Sampinute COVID-19 Antigen MIA | | SAMPINUTE™ Analyzer | BBB Inc. (distributed by Celltrion USA, Inc.) | SARS-CoV-2 receptor binding domains (RBDs) spike proteins | see further product documentation | see IFU | see IFU | magnetic force-assisted electrochemical sandwich immunoassay | US FDA EUA |
| 256091 256113 256114 | Bd Kit For Rapid Detection of SARS-CoV-2 | | see IFU | Becton, Dickinson and Company | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | Health Canada |
| 256082 | BD Veritor System for Rapid Detection of SARS-CoV-2 | 30T/kit | BD Veritor™ Plus Analyzer | Becton, Dickinson and Company | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | chromatographic digital immunoassay | US FDA EUA Health Canada PMDA TGA |
| | Wantai SARS-CoV-2 Ag Rapid Test (Colloidal Gold) | | | Beijing Wantai Biologicalpharmacy Enterprise Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| | SARS-CoV-2 Virus Antigen Detection Kit (Colloidal Gold Method) - POCT | | | BGI Europe A/S (Denmark) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| | SARS-CoV-2 Antigen Rapid Test (Colloidal Gold Method) | | | BIOHIT HealthCare (Hefei) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| | Istatis COVID-19 Antigen Test | | | Biolytical Laboratories Inc. (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 6 months | see IFU | see IFU | Visual read | Health Canada |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|-------------------------------------|------------------------------|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| RG1901DG | NowCheck COVID-19 Antigen Test | | | BioNote Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| COV9712A | C-Sync COVID-19 Antigen Test | 20 | n/a | Biosynchronicity Corporation | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | US FDA EUA |
| SW40006 | BIOSYNEX COVID-19 Ag BSS | see IFU | n/a | Biosynex Swiss SA | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | ANSM |
| SW40010 SW40010F | BIOSYNEX COVID-19 Ag+ | see IFU | n/a | Biosynex Swiss SA | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | ANSM |
| 2070088301 | Rapid Response™ COVID-19 Antigen Rapid Test Device | | n/a | BTNX Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | Health Canada |
| 2070089201 | Rapid Response™ COVID-19 Antigen Rapid Test Device | | n/a | BTNX Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | Health Canada |
| SWB-19 | Rapid Response™ COVID-19 Antigen Rapid Test Device | | n/a | BTNX Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | Health Canada |
| Rapid Response COV-19C25 | Rapid Response™ COVID-19 Antigen Rapid Test Device | | n/a | BTNX Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA Health Canada |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Surescreen Diagnostics COVID-19 Antigen Rapid Test Cassette | | n/a | BTNX Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| | SARS Coronavirus Antigen Kit Rapiim SARS-CoV-2-N PRT-C2N01A | | | Canon Medical Systems Corporation | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | | PMDA |
| | Quampas COVID-19 Antigen Test Kit | | | Cellspect Co.,Ltd. | | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | Aria® COVID-19 Ag Rapid Test | | n/a | CTK Biotech Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| R0182C | OnSite® COVID-19 Ag Rapid Test | 20Test/kit | n/a | CTK Biotech Inc | SARS-CoV-2 antigen | see WHO EUL Public Report | see IFU | nasal | Visual Read | TGA WHO EUL |
| R0182C | OnSite® COVID-19 Ag Rapid Test | 20Test/kit | n/a | CTK Biotech Inc | SARS-CoV-2 antigen | see further product documentation | see IFU | nasal | Visual Read | ANSM |
| | QuickNavi-COVID19 Ag | | | Denka Co., Ltd. | | see further product documentation | see IFU | see IFU | | PMDA |
| 311500 | LIAISON® SARS-CoV-2 Ag | see IFU | see IFU | DiaSorin | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Reader required | US FDA EUA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Exdia EK test COVID-19 Ag | | n/a | Eiken Chemical Co., Ltd. | SARS-CoV-2 antigen | see further product documentation | see IFU | nasal | Visual Read | PMDA |
| | Ellume COVID-19 Home Test | | see IFU | Ellume Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Reader required | US FDA EUA |
| ATG 900-207 ATG 900-208 ATG 900-210 | CovClear COVID-19 Antigen Test | | | Empowered Diagnostics LLC (United States Of America) | | 12 | | | | TGA |
| | Fuji Dry Chem IMMUNO AG Handy COVID-19 Ag | | | Fujifilm Corporation | | see further product documentation | see IFU | see IFU | | PMDA |
| | Accuraseed SARS-CoV-2Ag | | | FUJIFILM Wako Pure Chemical Corporation | | see further product documentation | see IFU | see IFU | | PMDA |
| | TEGARUNA Stick SARS-CoV-2 Ag | | | Fujinaga Pharmaceutical- Co., Ltd | | see further product documentation | see IFU | see IFU | | PMDA |
| | Lumipulse SARS-CoV-2 Ag | | | Fujirebio Inc | | see further product documentation | see IFU | see IFU | | PMDA |
| | Lumipulse Presto SARS-CoV-2 Ag | | | Fujirebio Inc | | see further product documentation | see IFU | see IFU | | PMDA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| 231906 | ESPLINE SARS-CoV-2 | 100T/kit | n/a | Fujirebio Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | PMDA |
| COVAG025-U COVAG025-NU | GenBody COVID-19 Ag | 25T/kit | | GenBody Inc (Korea - Republic of) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 21 Months | see IFU | Visual read | US FDA EUA TGA |
| | 2019-nCoV Ag Saliva Rapid Test Card | | | Guangzhou Decheng Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| W634P0013 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 20T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | WHO EUL |
| W634P0014 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 100T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | WHO EUL |
| W634P0015 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 20T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | WHO EUL |
| W196P0019 W196P0020 W196P0021 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 1T/kit 5T/kit 20T/kit | | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| W196P0031 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | see IFU | Visual read | Health Canada |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | GSD NovaGen SARS CoV-2 Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| INCP-G502 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | Health Canada |
| ICOV-802 | COVID-19 Antigen Rapid Test (Oral Fluid) ICOV-802 | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| ICOV-502 | COVID-19 Antigen Rapid Test (swab) ICOV-502 | 25 tests/kit | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | Nasopharyngeal | | TGA |
| | COVID-19 Antigen Rapid Test Cassette | | | HANGZHOU BIOTEST BIOTECH NO LTD (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| | Clungene Covid-19 Antigen Rapid Test | | | Hangzhou Clongene Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| | LYHER Novel Coronavirus (Covid-19) Antigen Test Kit (Colloidal Gold) | | | Hangzhou Laihe Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| 303035-25 303035-40 | Novel Coronavirus (COVID-19) Antigen Test (Colloidal Gold) | 25 T/kit 40 T/kit | | Hangzhou Laihe Biotech Co., Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | WHO EUL |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|---|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Cellife Covid-19 Antigen Test Cassette | | | Hangzhou Testsea Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| | Testsea SARS-CoV-2 Antigen Test Kit | | | Hangzhou Testsea Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| CT-P60 D-2 02 | Celltrion DiaTrust™ COVID-19 Ag Rapid Test | 25T/kit | n/a | Humasis, Co Ltd. (distributed by Celltrion USA, Inc) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 24 months | see IFU | Visual Read | US FDA EUA |
| ICO-3000P | iHealth COVID-19 Antigen Rapid Test Pro | 40T/kit | n/a | iHealth Labs, Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 15 months | nasal | Visual read | US FDA EUA |
| | GoToKnow COVID-19 Antigen Rapid Test | 40T/kit | n/a | iHealth Labs, Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| COVAG-RC | SCoV-2 Ag Detect Rapid Test | 50T/kit | n/a | InBios International, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| | InnoScreen COVID-19 Antigen Rapid Test Device | | | Innovation Scientific Pty Ltd (Australia) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| | 2019 nCoV Ag Test (Latex Chromatography Assay) – POCT | | n/a | Innovita (Tangshan) Biological Technology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| | PixoTest® COVID-19 AG Test Kit | | | iXensor Co Ltd (Taiwan) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| | SARS-CoV-2 antigen Test Kit (LFIA) | | n/a | Jiangsu Medomics medical technology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | TGA |
| | KANEKA Immunochromatography SARS-CoV-2 Ag | | n/a | KANEKA CORPORATION | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | PMDA |
| | KBM line check nCoV (stick type) | | n/a | Kojin Bio Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | PMDA |
| | KBM LineCheck nCoV/Flu | | n/a | Kojin Bio Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | PMDA |
| | LUCA NK COVID-19 Ag NP LLB | | | LUCA AICell Inc Republic of Korea | | | | Nasopharyngeal, nasal, and saliva sample | | TGA |
| | LUCA NK COVID-19 Ag nasal LLB | | | LUCA AICell Inc Republic of Korea | | | | Nasopharyngeal, Nasal and saliva sample | | TGA |
| | LUCA NK COVID-19 Ag Saliva LLB | | | LUCA AICell Inc Republic of Korea | | | | Nasopharyngeal, Nasal and saliva sample | | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| Lo16000110012 Lo16000110024 Lo16000110048 | LumiraDx SARS-CoV-2 Ag Test | 12T/kit 25T/kit 48T/kit | LumiraDx Platform | LumiraDx UK Ltd. | SARS-CoV-2 nucleocapsid protein antigen | 6 | see IFU | see IFU | Reader required | US FDA EUA WHO EUL |
| Lo16000113012 Lo16000113024 Lo16000113048 | LumiraDx SARS-CoV-2 Ag Test | 12T/kit 25T/kit 48T/kit | LumiraDx Platform | LumiraDx UK Ltd. | SARS-CoV-2 nucleocapsid protein antigen | 10 months | see IFU | see IFU | Reader required | Health Canada |
| CP0023 | COVIDx-SARS-CoV-2 Rapid Antigen Test Kit | 25T/kit | LumiraDx Platform | Lumos Diagnostics Ata Rapid Pathogen Screening Inc Ata Lumos Diagnostics Inc (United States) | SARS-CoV-2 nucleocapsid protein antigen | 12 | see IFU | see IFU | | Health Canada |
| | STANDARD Q COVID-19 Ag Test | | | MALCOM COMPANY LIMITED | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | GLINE-2019-nCoV Ag Kit | | | Medical & Biological Laboratories, Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | Inspector Kowa SARS-CoV-2 | | | Medical & Biological Laboratories, Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| MS-TRA-5 | Medsup COVID-19 Rapid Antigen Test | | n/a | Medsup Medical (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 12 | see IFU | see IFU | | Health Canada |
| | Fuji Dry Chem IMMNO AG Cartridge COVID-19 Ag | | n/a | Mizuhomedy Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | PMDA |

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| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Quick Chaser Auto SARS-CoV-2 | | n/a | Mizuhomedy Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Reader required | PMDA |
| | Rapid SARS-CoV-2 Antigen Test Card | | | MP Biomedicals Asia Pacific Pte Ltd (Singapore) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | | | | TGA |
| ND-MD8147 | Nano-Check COVID-19 Antigen Test | 20T/kit | n/a | Nano-Ditech Corp | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Immunofine SARS-COV-2 | | | Nichirei Biosciences Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| 828-100 | ASSURE-100 Rapid COVID-19 Test | 30T/kit | n/a | Oceanit Foundry LLC | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| 1001-0614 1001-0615 | InteliSwab COVID-19 Rapid Test Pro | 25T/kit 100T/kit | n/a | OraSure Technologies, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | see IFU | Visual read | US FDA EUA |
| 1001-0614.001 | InteliSwab COVID-19 Rapid Test Pro | | n/a | OraSure Technologies, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | see IFU | Visual read | Health Canada Interim Order |
| 619 9949 | VITROS Immunodiagnostic Products SARS-CoV-2 Antigen Reagent Pack | 100T/kit | VITROS 3600 Immunodiagnostic System and the VITROS 5600/XT 7600 Integrated Systems | Ortho Clinical Diagnostics, Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Analyzer required | US FDA EUA Health Canada PMDA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|--|---|---|-----------------------------------|---------------------------------|----------------|-------------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| COV05 | Pcl COVID19 AG Rapid Fia | | Pcl Immunofluorescence Analyzer Pclok Ez | Pcl Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Analyzer required | TGA |
| COV04S-1T COV04S-2T | Pcl COVID19 AG Gold | 1T/kit 2T/kit | n/a | Pcl Inc. (South Korea) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | Health Canada |
| | INDICAID COVID-19 Rapid Antigen Test | 25T/kit | n/a | PHASE Scientific International Limited | SARS-CoV-2 nucleocapsid protein antigen | 15 months | see IFU | see IFU | Visual read | US FDA EUA |
| P0092 P0093 P0094 P0095 | INDICAID COVID-19 Rapid Antigen Test | 25T/kit | n/a | PHASE Scientific International Limited | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | see IFU | Visual read | Health Canada |
| SS03P25 | Sure Status COVID-19 Antigen Card Test | 25T/kit | n/a | Premier Medical Corporation Private Limited | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | Nasopharyngeal | Visual read | WHO EUL |
| SS03-NS-P25 | Sure Status COVID-19 Antigen Card Test | 25T/kit | n/a | Premier Medical Corporation Private Limited | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | nasal | Visual read | WHO EUL |
| 09COV30D | SARS-CoV-2 Antigen Rapid Test (without software) | | n/a | Qingdao Hightop Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| 20387 | QuickVue SARS Antigen Test | 25T/kit | n/a | Quidel Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|-------------------------------|-------------------------|---|--|---|-----------------------------------|---------------------------------|---------------|-----------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| 20396 | QuickVue SARS Antigen Test | 25T/kit | n/a | Quidel Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | Health Canada |
| 20374 | Sofia SARS Antigen FIA | 25T/kit | Sofia and Sofia 2 instrument | Quidel Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Reader required | US FDA EUA TGA |
| QPR8302 | Omnia SARS-CoV-2 Antigen Test | 20T/kit | Qorvo Biotechnologies Omnia System (Catalog number # QPR9002) | Qorvo Biotechnologies, LLC | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Reader required | US FDA EUA |
| | BIOCREDIT COVID-19 Ag | | | RapiGEN | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | see IFU | TGA PMDA |
| G61RHA20 G61RHA20C | BIOCREDIT COVID-19 Ag | | | RapiGEN | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | see IFU | WHO EUL |
| C1120-25 C1120-100 | Ramp COVID-19 Antigen Test | 25T/kit 100T/kit | | Response Biomedical Corporation (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | see IFU | Reader required | Health Canada |
| | Eclusis Reagent SARS-CoV-2 Ag | | n/a | Roche Diagnostics Co., Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | see IFU | PMDA |
| | SARS-Cov-2 Rapid Antigen Test | | n/a | Roche Diagnostics Co., Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | PMDA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|---|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | SARS-Cov-2 Rapid Antigen Test II | | n/a | Roche Diagnostics Co., Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | PMDA |
| CLA-COV19AG-VIS/102241 | Sienna-Clarity COVID-19 Antigen Rapid Test Cassette | | n/a | Salofa Oy | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 21 Months | see IFU | Visual Read | US FDA EUA |
| | Clarity COVID-19 Antigen Rapid Test Cassette | | n/a | Salofa Oy | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | US FDA EUA |
| | OVIOS COVID-19 Antigen Rapid Test Cassette | | n/a | Salofa Oy | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | US FDA EUA |
| | Spring Health COVID-19 Antigen Rapid Test Cassette | | n/a | Salofa Oy | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | US FDA EUA |
| | Salocor COVID-19 Antigen Rapid Test Cassette | | n/a | Salofa Oy | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | US FDA EUA |
| | CO-Check COVID-19 Rapid Antigen Test (POCT) | | n/a | Sanwa BioTech Limited (Hong Kong - SAR of China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| | CO-Check SARS-CoV-2 Antigen LFIA Test (POCT) | | n/a | Sanwa BioTech Limited (Hong Kong - SAR of China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|---|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|--|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | ALiA SARS-CoV-2 Antigen FIA Test (POCT) | | n/a | Sanwa BioTech Limited (Hong Kong - SAR of China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| | STANDARD Q COVID-19 Ag Test 2.0 | | n/a | SD Biosensor, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | USFDA EUA |
| 09COV30D | STANDARD Q COVID-19 Ag Test | | n/a | SD Biosensor Inc. | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | see IFU | Visual Read | TGA WHO EUL |
| 9901-NCOV-01G | SARS-Cov-2 Rapid Antigen Test | | n/a | SD Biosensor Inc. (distributed by Roche Diagnostics Australia) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| 9901-NCOV-04G | SARS-Cov-2 Rapid Antigen Test | | n/a | SD Biosensor Inc. (distributed by Roche Diagnostics Australia) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | Health Canada |
| 9901-NCOV-06G | SARS-Cov-2 Rapid Antigen Test | | n/a | SD Biosensor Inc. (distributed by Roche Diagnostics Australia) | SARS-CoV-2 nucleocapsid protein antigen | 24 | see IFU | see IFU | Visual Read | Health Canada |
| 09COV30D | STANDARD™ Q COVID-19 Ag Test Nasal Swab | | n/a | SD Biosensor Inc. (distributed by Bioelect Pty Ltd) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual Read | TGA |
| | RapidTesta SARS-CoV-2 | | n/a | SEKISUI MEDICAL CO., LTD. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|--------------------------|--|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| R-425-25-T | Diagnostic Kit for COVID-19 Antigen Test (Colloidal Gold) | 25Test/kit | n/a | Shanghai Kehua Bioengineering Co. Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see WHO EUL Public Report | see IFU | see IFU | Visual Read | WHO EUL |
| | Clinitest COVID-19 Antigen Rapid Test | | n/a | Siemens Healthcare Diagnostics Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | PMDA |
| | SARS-CoV-2 Antigen(CoV2Ag) | | n/a | Siemens Healthcare Diagnostics Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | SGTi-flex COVID-19 Ag | | | Sugentech Inc Republic of Korea | | | | Nasopharyngeal and Nasal | | TGA |
| | SureScreen SARS-CoV-2 Antigen Rapid Test Cassette (Nasal Swab) (Gold) for professional use | | n/a | SureScreen Diagnostics Ltd (United Kingdom) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | SARS-CoV-2 Antigen Rapid Test Cassette (Rhino Swabs) (POCT) | | n/a | SureScreen Diagnostics Ltd (United Kingdom) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | HISCL SARS-CoV-2 Ag Assay Kit | | | Sysmex Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | | PMDA |
| | AFIAS COVID-19 Ag Test Cartridge | | | TOKYO BOEKI MEDISYS INC | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | CL AIA-PACK SARS-CoV-2-Ag | | | TOSOH CORPORATION | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|---|--|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|--|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | HEALGEN COVID-19 Antigen rapid test | | | Takara Bio Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | Immuno Ace SARS-CoV-2 / Capilia SARS-CoV-2 | | | TAUNS LABORATORIES, INC. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | ImmunoArrow SARS-CoV-2 | | | TOYOBO CO., LTD. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| 1N40C5-1-US | Rapid SARS-CoV-2 Antigen Test Card | 1T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-2-US | Rapid SARS-CoV-2 Antigen Test Card | 2T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-4-US | Rapid SARS-CoV-2 Antigen Test Card | 4T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-5-US | Rapid SARS-CoV-2 Antigen Test Card | 5T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-8-US | Rapid SARS-CoV-2 Antigen Test Card | 8T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-10-US | Rapid SARS-CoV-2 Antigen Test Card | 10T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| 1N40C5-20-US | Rapid SARS-CoV-2 Antigen Test Card | 20T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|------------------------------------|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| 1N40C5-40-US | Rapid SARS-CoV-2 Antigen Test Card | 40T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | 11 months | see IFU | Visual read | US FDA EUA |
| XH-100-110 | SPERA COVID-19 Ag Test | 10T/kit | n/a | Xtrava Health | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| N/A- NOT APPLICABLE | | | | | | | | | | |
| Disclaimer:The Global Fund does not endorse or warrant the fitness of any product on the List for a particular purpose. In addition, the Global Fund assumes no responsibility for any misstatement or omission from the list and directs Principal Recipients of Global Fund grants to conduct their own independent confirmation that the information on a given product on the list is accurate before relying on it to make a purchase order for that product, and to ensure that any purchase is in compliance with all the requirements of the Global Fund’s quality assurance policy. The Global Fund does not warrant or represent that the products listed have obtained regulatory approval for use in any particular country of the world, or that their use is otherwise in accordance with the national laws and regulations of any country, including, but not limited to, intellectual property laws. The Global Fund disclaims any and all liability and responsibility for any injury, death, damage or loss of any kind whatsoever that may arise as a result of, or in connection with the procurement, distribution and use of any product included in the list. | | | | | | | | | | |

List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)

The following emergency procedures established by WHO and the Regulatory Authorities of the Founding Members of the GHTF have been identified by the QA Team and will be used to determine eligibility for procurement of COVID-19 diagnostics. The product, to be considered as eligible for procurement with GF resources, shall be listed in one of the below mentioned lists:

- WHO Prequalification decisions made as per the Emergency Use Listing (EUL) procedure opened to candidate in vitro diagnostics (IVDs) to detect SARS-CoV-2;
- The United States Food and Drug Administration’s (USFDA) general recommendations and procedures applicable to the authorization of the emergency use of certain medical products under sections 564, 564A, and 564B of the Federal Food, Drug, and Cosmetic Act;
- The decisions taken based on the Canada’s Minister of Health interim order (IO) to expedite the review of these medical devices, including test kits used to diagnose COVID-19;
- The COVID-19 diagnostic tests approved by the Therapeutic Goods Administration (TGA) for inclusion on the Australian Register of Therapeutic Goods (ARTG) on the basis of the Expedited TGA assessment
- The COVID-19 diagnostic tests approved by the Ministry of Health, Labour and Welfare after March 2020 with prior scientific review by the PMDA
- The COVID-19 diagnostic tests listed on the French government website and under the control of the French Health Authority ANSM

The following websites provide access to Instructions For Use of certain products:

- <https://extranet.who.int/pqweb/vitro-diagnostics/coronavirus-disease-covid-19-pandemic-%E2%80%94-emergency-use-listing-procedure-eul-open>
- <https://www.fda.gov/medical-devices/emergency-situations-medical-devices/emergency-use-authorizations>

The following website provides WHO Interim Guidance “Diagnostic testing for SARS-CoV-2”:
<https://www.who.int/publications/i/item/diagnostic-testing-for-sars-cov-2>

Important Note: The following lists are not exhaustive.

SARS-CoV-2 Nucleic Acid Amplification Technologies

| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
|---------------------------------------|---------------------------|------------------|---|--|--|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| 444213 | 1COPY COVID-19 QPCR KIT | 100T/kit | QIAamp Viral RNA Mini Kit RNA mini kit (QIAgen) | Light Cycler 480 (Roche) | 1DROP INC. (imported by Luminarie Canada Inc.) | E gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Rotor-Gene Q 5plex HRM (Qiagen) | | | | | | | |
| | | | | Applied Biosystems Quantstudio5 (Thermo Fisher Scientific) | | | | | | | |
| | | | | Applied Biosystems 7500 Real-Time PCR Instrument system (Thermo Fisher Scientific) | | | | | | | |
| | | | | CFX96™ Real-Time PCR Detection system (BIO-RAD) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------------------|------------------|--|--|---|------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 3B304 | TRUPCR SARS-CoV-2 Kit | 100T/kit | TRUPCR® Viral RNA Extraction Kit (3B213V/3B214V) | Applied Biosystems Quantstudio3 (Thermo Fisher Scientific) | 3B Blackbio Biotech India Ltd (a Kilpest India Ltd company) | RdRp, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Rotor-Gene Q 5plex HRM (Qiagen) | | | | | | | |
| 3103010011 | 3DMed 2019-nCoV RT-qPCR Detection Kit | 100T/kit | ANDiS Viral RNA Auto Extraction & Purification Kit | Applied Biosystems 7500 Real-Time PCR Instrument system (Thermo Fisher Scientific) | 3D Biomedicine Science & Technology Co., Ltd. | N, E and ORF-1ab genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | Automated Nucleic Acid Extraction System ANDiS 350 | | | | | | | | |
| | | | Qiagen DSP Viral RNA Mini Kit | | | | | | | | |
| | 3EO Health COVID-19 Test | | | | 3EO Health, Inc | see IFU | See IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 190-000 | ID NOW COVID-19 Test Kit | 96T/kit | ID NOW Instrument | | Abbott Diagnostics Scarborough Inc | RdRp segment | 24 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order PMDA TGA |
| 193-000 | ID NOW COVID-19 2.0 Test Kit | 96T/kit | ID NOW Instrument | | Abbott Diagnostics Scarborough Inc | RdRp segment | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| 09N78-090 | Alinity m SARS-CoV-2 AMP Kit | 192T/kit | Alinity m System | | Abbott Molecular | RdRp and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| 09N78-095 | Alinity m SARS-CoV-2 AMP Kit | 96T/kit | Alinity m System | | Abbott Molecular | RdRp and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA PMDA TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------------------|-----------------------------------|--|---|---|--------------------------------|---------------------------------|---------------------------------|---------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 09N77-090 | Abbott RealTime SARS-CoV-2 RT-PCR Kit | 96T/kit | Abbott m2000 | | Abbott Molecular | RdRp and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order TGA WHO EUL |
| 09N77-095 | Abbott RealTime SARS-CoV-2 RT-PCR Kit | 96T/kit | Abbott m2000 | | Abbott Molecular | RdRp and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Resolute 2.0 SARS CoV-2 Detection Kit | see IFU | see IFU | | Accelerate Technologies Pte Ltd (DxD Hub) | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| 13279F 13278D 13281D | MassARRAY® SARS-CoV-2 Panel | 960T/kit 3840T/kit 768T/kit | NucliSENS® easyMAG® (bioMérieux) | MassARRAY System | Agena Bioscience, Inc. | N gene, ORF-1 and ORF-1ab gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | KingFisher Flex Purification System (ThermoFisher) | | | | | | | | |
| 821025 | RealStar® SARS-CoV-2 RT-PCR Kit U.S. | 384T/kit | AltoStar® Automation System AM16 | CFX96™ Touch Real-Time PCR Detection System (Bio-Rad) | Altona Diagnostics GmbH | E and S gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96™ Touch Deep Well Real-Time PCR Detection System (BioRad | | | | | | | |
| | RESOLUTE 2.0 | | | | AMT Pte Ltd (Singapore) | | | | | | TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|--------------------------|----------------|---------------------------------|---------------------------------|---------------|---|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 64-Co304 | BioCode® SARS-CoV-2 Assay | 384T/kit | NucliSENS® easyMAG® (bioMérieux) | BioCode® MDx-3000 automated system | Applied BioCode Inc | N gene | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA |
| | | | MagNA Pure 96 (Roche) | | | | | | | | |
| DX-1001-001-000 | Linea™ COVID-19 Real-Time PCR Assay Kit assay kit | 100T/kit | QIAamp® Viral RNA Mini Kit (Qiagen) | Applied Biosystems QuantStudio™ Dx Real-Time PCR system | Applied DNA Sciences | S gene | 6 months | see IFU | see IFU | Under risk assessment for Omicron variant Note: See WHO Information Notice for IVD users 2021/01 with regards to mutations in SARS-CoV-2 | US FDA EUA |
| DX-1001-002-000 | | 500T/kit | TRIzol™ RNA Extraction Kit (Invitrogen) | Applied Biosystems QuantStudio 5 Real-Time PCR System | | | | | | | |
| DX-1001-003-000 | | | 1000T/kit | Mag-Bind Viral RNA Xpress Kit (Omega Bio-Tek) automated on the Hamilton STARlet system | | | | | | | |
| | Identity Pack SARS-CoV-2 | | | | Arkray Factory Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|--|--|---|-----------------------------|---------------------------------|---------------------------------|---------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| iAMP-COVID19-100 | iAMP® COVID-19 Detection Kit | 100T/kit | not required | CFX96 Real-Time System (Bio-Rad) | Atila BioSystems, Inc. | N gene and the ORF-1ab gene | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Roche LightCycler 480 Instrument II Real-Time PCR System | | | | | | | |
| | | | | Atila PowerGene 9600 Plus Real-Time PCR System | | | | | | | |
| | | | | Applied Biosystems 7500 Fast Real-Time PCR System | | | | | | | |
| 445003-01 | BD SARS-CoV-2 Reagents | 24T/kit | BD MAX™ System | | Becton, Dickinson and Company | N gene (N1 and N2 regions) | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA Health Canada/Interim Order |
| MFG030010 | Real-Time Fluorescent RT-PCR Kit for Detecting SARS-2019-nCoV | 50T/kit | TIANamp Virus RNA extraction Kit (DP315-R) TIANGEN | Applied Biosystems 7500/7500 Fast Real-Time PCR System | BGI Europe A/S | ORF1ab | 6 months | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | QIAamp Virus RNA Mini Kit (cat. #52904). | Applied Biosystems QuantStudio 5 Real-Time PCR Systems | | | | | | | |
| | | | RNA extraction kit by MGI Tech (Wuhan) (No.20200167) | SLAN-96P PCR system | | | | | | | |
| | | | | LightCycler® 480 System | | | | | | | |
| | Real-Time Fluorescent RT-PCR Kit for Detecting SARS-Cov-2 (2 gene) | | | | BGI Europe A/S (distributed by BGI Health (AU) Company Pty Ltd) | | | | | | TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|--|----------------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| MFG030010 | Real-Time Fluorescent RT-PCR Kit for Detecting SARS-2019-nCoV | 50T/kit | QIAamp Virus RNA Mini Kit (cat. #52904 or 52906). | Applied Biosystems 7500 Real-Time PCR System | BGI Genomics Co. Ltd | ORF1ab | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| CT8233-48T | Multiple Real-Time PCR Kit for Detection of 2019-nCoV | 48T/kit | Nucleic acid extraction Kit (XABT, Cat. # CN8033) | Applied Biosystems 7500 Real-Time PCR System | Beijing Applied Biological Technologies Co. Ltd., (XABT) | ORF1ab and N gene and the E gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | QIAamp Viral RNA Mini Kit (Qiagen, Cat. # 52904 or 52906) | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | PURELINK VIRAL RNA/DNA KIT (Invitrogen, Cat. #12280050) | LightCycler 480 (System II) (Roche) | | | | | | | |
| | | | High Pure Viral RNA Kit (Roche, Cat. # 11858882001) | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|-------------------------------------|----------------------|---|--|--|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| WS-1248 | Wantai SARS-CoV-2 RT-PCR Kit | 48T/kit | Beijing Wantai Nucleic Acid Extraction Kit (cat. # ZCT1246) | Applied Biosystem® 7500 Real-Time PCR system | Beijing Wantai Biological Pharmacy Enterprise Co., Ltd | ORF1ab and N gene | see IFU | see IFU | see IFU | See IFU | WHO EUL |
| | | | Beijing Wantai Nucleic Acid Extraction Kit on KingFisher Flex 96 (ThermoFisher) | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | Beijing Wantai Nucleic Acid Extraction Kit on NEXOR 32 (Yantai Addcare Bio-Tec) | | | | | | | | |
| | | | Beijing Wantai Nucleic Acid Extraction Kit on NEXOR 96 (Yantai Addcare Bio-Tec) | | | | | | | | |
| | | | QIAamp Viral RNA Mini Kit QIAGEN (cat. # 52094) | | | | | | | | |
| BC-01-0099 BC-01-0099 x4 | BioCore 2019-nCoV Real Time PCR Kit | 100T/kit 400T/kit | QIAamp DSP Viral RNA Mini Kit (Qiagen; catalog #61904) | Applied Biosystems 7500 Real-Time PCR System | BioCore Co. Ltd. | N gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA TGA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | SLAN-96P (Shanghai Hongshi Medical Technology Co. Ltd) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|-----------------------|---|--|--|-----------------|---------------------------------|---------------------------------|----------------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| BS-SY-SC2-100 BS-SY-SC2-1000 | Bio-Speedy® Direct RT-qPCR SARS-CoV-2 | 100T/kit 1000T/kit | | LightCycler 96 (Roche) | Bioeksen R&D Technologies Ltd | ORF1ab gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | Rotor-Gene Q (Qiagen) | | | | | | | |
| BS-SY-SC2-100 BS-SY-SC2-1000 | Bio-Speedy® Direct RT-qPCR SARS-CoV-2 rebranded to BioeXsen SARS-CoV-2 RT PCR | 100T/kit 1000T/kit | | LightCycler 96 (Roche) | Bioeksen R&D Technologies Ltd (distributed by BioeXsen GmbH) | ORF1ab gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | Rotor-Gene Q (Qiagen) | | | | | | | |
| BS-SY-WCOR-304-100 | Bio-Speedy® Direct RT-qPCR SARS-CoV-2 | 100T/kit | RINA M14 Nucleic Acid Extraction Robot (Cat No: RINA-M14-01) | LightCycler 96 (Roche) | Bioeksen R&D Technologies Ltd | ORF1ab gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | Rotor-Gene Q 5plex (Qiagen) | | | | | | | |
| 423745 | BioFire® COVID-19 Test | 6T/kit | FilmArray® 2.0 and/or the FilmArray® Torch Instrument Systems | | BioFire Defense, LLC | ORF1ab and ORF8 | see IFU | see IFU | nasopharyngeal swabs | For consumables and details of componants refer to IFU | US FDA EUA |
| 423744 | | 30T/kit | | | | | | | | | |
| 423738 | Biofire Respiratory Panel 2.1 (RP2.1) | | FilmArray® 2.0 and/or the FilmArray® Torch Instrument Systems | | BioFire Diagnostics LLC | see IFU | 18 months | see IFU | see IFU | For consumables and details of componants refer to IFU | (US FDA EUA replaced by DeNovo approval) Health Canada/Interim Order TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|-------------------------------------|------------------|---|--|--------------|-------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 500-003-XMP | BioGX Xfree COVID-19 Direct RT-PCR | 104T/kit | n/a | Applied Biosystems QuantStudio 5 | BioGX, Inc. | N gene | 15 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 Fast Dx | | | | | | | |
| | | | | BioGX pixl.16 platform | | | | | | | |
| | | | | Bio Molecular Systems Magnetic Induction Cyclers (MIC) | | | | | | | |
| | | | | Bio-Rad CFX396 Touch | | | | | | | |
| | | | | Bio-Rad CFX384 Touch | | | | | | | |
| | Biomeme SARS-CoV-2 Real-Time RT-PCR | 96T/kit | MagMax Viral/Pathogen on KingFisher Flex Purification System (ThermoFisher) | Bio-Rad CFX96 | Biomeme Inc. | ORF1ab and S gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | Biomeme M1 Sample Prep Cartridge Kit for RNA 2.0 | QuantStudio 5 | | | | | | | |
| 3000555 | Biomeme SARS-CoV-2 Go-Strips | | Biomeme’s Franklin Real-Time PCR System | | Biomeme Inc. | ORF1ab and S gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA Health Canada/Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|------------------------------|------------------------------------|---|--|--------------------------|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 424017 | ARGENE® COVID-19 R-GENE® | 30 and 120T/kit (depending on PCR) | EMAG® (bioMérieux) | 7500 & 7500 Fast Real-Time PCR System (Applied Biosystems) | BioMérieux SA | N gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | ANSM |
| | | | | QuantStudio 5 and 5 DX (Applied Biosystems) | | | | | | | |
| | | | NucliSENS easyMAG (bioMérieux) | LightCycler 480 (System II) (Roche) | | | | | | | |
| | | | MGISP-960 | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | QIASymphony SP (QIAgen) | CFX Opus 96 | | | | | | | |
| | | | MagNA Pure 96 (Roche) | Rotor-Gene Q (Qiagen) | | | | | | | |
| | SARS-COV-2 R-GENE® | | | | BioMérieux SA | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| 12013743 | Bio-Rad SARS-CoV-2 ddPCR Kit | 200T/kit | ThermoFisher MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (Cat No. A48310, 1000 reactions) | QX200™ PCR Systems | Bio-Rad Laboratories Inc | P and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | QIAamp Virus RNA Mini Kit (cat. # 52906) | | | | | | | | |

SARS-CoV-2 Nucleic Acid Amplification Technologies

| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
|---------------------------------------|--------------------------------|------------------|---|---|--------------|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| | | | ThermoFisher MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (Cat No. A48310, 1000 reactions) on KingFisher Flex system | QXDx™ Droplet Digital™ PCR Systems | | | | | | | |
| BS7nCoV | Real-Q 2019-nCoV Detection Kit | 100T/kit | MagNA Pure 96 (Roche) or manual | 7500 Real-Time PCR System (Applied Biosystems) | BioSewoom | E gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | 7500 Fast Real-Time PCR System (Applied Biosystems) | | | | | | | |
| | | | | QuantStudio5 realtime PCR instrument (Applied Biosystems) | | | | | | | |
| | | | | CFX96 DX Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | CFX96 Real-Time PCR Detection System (Bio-Rad) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--------------------------------|------------------|--|---|--------------|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 3150058 | BIOSYNEX AMPLIQUICK SARS-CoV-2 | 96T/kit | QIAamp Mini RNA Viral Extraction Kit (QIAGEN) | QuantStudio5 realtime PCR instrument (Applied Biosystems) | Biosynex | E gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | ANSM |
| | | | BIOSYNEX AMPLIQUICK® Lysis | 7500 Real-Time PCR System (Applied Biosystems) | | | | | | | |
| | | | Liferiver Viral Extraction Kit (Ref: ME- 0044) | LightCycler 480 (System II) (Roche) | | | | | | | |
| | | | | DT lite 48/96 (DNA Technology / Amplix) | | | | | | | |
| | | | | QuantGene 9600 (BIOER) | | | | | | | |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | SLAN-48/96P (Sansure) | | | | | | | |
| | | | | LineGene Mini S (Bioer) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------|------------------|---|--|--------------|-----------------------------|---------------------------------|---------------------------------|-----------------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| TD1100 | COVID-19 RT-PCR PNA kit | 24T/kit | RNeasy Mini kit (Qiagen) | 7500 Real-Time PCR System (Applied Biosystems) | BioTNS | RdRp gene and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | 7500 Fast Real-Time PCR System (Applied Biosystems) | | | | | | | |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| XP3SARS-COV2-10 | Xpert® Xpress SARS-CoV-2 | 10T/kit | GeneXpert Xpress System (Tablet and Hub Configurations) | | Cepheid | RdRp gene and N and E genes | see IFU (302-7069) | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| XP3SARS-COV2-10 | Xpert® Xpress CoV-2 plus | 10T/kit | GeneXpert Dx with 6- and 10-color optical modules) | | Cepheid | RdRp gene and N and E genes | see IFU (302-7070) | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | GeneXpert Infinity Systems | | | | | | | | |
| XPRSARS-COV2-10 | Xpert® Xpress SARS-CoV-2 | 10T/kit | GeneXpert Xpress System (Tablet and Hub Configurations) | | Cepheid | Genes N2 and E | see IFU (302-3750) | see IFU | see IFU (302-3750) | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order TGA |
| XPRSARS-COV2-10 | Xpert® Xpress SARS-CoV-2 | 10T/kit | GeneXpert Dx with 6- and 10-color optical modules) | | Cepheid | Genes N2 and E | see IFU (302-3562) | see IFU | see IFU (302-3562) | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order TGA |
| | | | GeneXpert Infinity Systems | | | | | | | | |
| XPRSARS-COV2-10 | Xpert® Xpress SARS-CoV-2 | 10T/kit | GeneXpert Dx with 6-color optical modules | | Cepheid | Genes N2 and E | see IFU (302-3562) | see IFU | see IFU (302-3562) | For consumables and details of componants refer to IFU | WHO EUL |
| | | | GeneXpert Infinity Systems | | | | | | | | |
| XP3SARS-COV2-10 | Xpert Xpress CoV-2 plus | 10 T/kit | GeneXpert Instrument System (Dx and Infininty systems) | | Cepheid | RdRp gene and N and E genes | 12 months | 2-28°C | nasophareng eal/nasal | For consumables and details of componants refer to IFU | WHO EUL |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|---|---|-------------------------------|--------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | VIASURE SARS-CoV-2 Real Time PCR Detection Kit | | Viasure RNA-DNA Extraction kit (VIASURE) | Applied Biosystems 7500 Fast Real-Time PCR System | CerTest Biotec SL / Abacus dx | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | | | Maxwell® 16 Viral Total Nucleic Acid Purification Kit, using the Maxwell® 16 instrument (Promega) | Applied Biosystems StepOne™ Real-Time PCR System | | | | | | | |
| | | | Total Nucleic Acid Isolation (TNAI) Kit, using COBAS® AmpliPrep (ROCHE) | Bio-Rad CFX96™ Real-Time PCR System | | | | | | | |
| | | | MagDEA Dx SV kit, using the magLEAD® 12gC instrument (Precision System Science Co.) | Agilent Technologies AriaMx Real-Time PCR System | | | | | | | |
| | | | MagCore® Viral Nucleic Acid Extraction kit, using the MagCore® HF16 automated Nucleic Acid Extractor System | DNA-Technology Detection DTprime Real-time Detection Thermal Cycler | | | | | | | |
| | | | EZ1 Virus Mini Kit, using EZ1 instrument (Qiagen) | DNA-Technology DTLite Real-Time PCR System | | | | | | | |
| | | | mSample Preparation Systems RNA, using the Abbott m2000 RealTime System (Abbott Molecular) | Rotor-Gene® Q (Qiagen) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|--------------------------------------|--------------------|---------------------------------|---------------------------------|----------------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | | | | SmartCycler (Cepheid) | | | | | | | |
| | | | | Roche Molecular Diagnostics Cobas z480 Analyzer | | | | | | | |
| | | | | VIASURE 48 Real Time PCR System | | | | | | | |
| | | | | VIASURE 96 Real Time PCR System | | | | | | | |
| | VIASURE SARS-CoV-2 S gene Real Time PCR Detection Kit | | | | CerTest Biotec SL / Becton Dickensen | | see IFU | see IFU | nasopharyngeal swabs | For consumables and details of components refer to IFU | TGA |
| HBRT-COVID-19 | COVID-19 Real-Time PCR Kit | 24T/kit | Thermo Scientific™ KingFisher™ Flex | 7500 Real-Time PCR System (Applied Biosystems) | Chaozhou HybriBio Biochemistry Ltd. | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | TGA WHO EUL |
| | | | Bioer GenePure Pro Nucleic Acid Purification System | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | | 96S Real-Time PCR System (SLAN) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|---|---------------------------------------|--|-----------------------------|---------------------------------|---------------------------------|----------------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| TR-US-01 | Clinomix TrioDx RT-PCR COVID-19 Test | 100T/kit | QIAamp Viral RNA Mini Kit | ABI QuantStudio 6 Flex | Clinomix USA Inc. | RdRp gene and N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | Maxwell RSC Viral Total Nucleic Acid Multi-Pack Kitwith the Maxwell RSC 48 instrument | | | | | | | | |
| | | | MagMAX Viral/Pathogen II Nucleic Acid Isolation Kit with KingFisher Flex instrument | | | | | | | | |
| COVID-K-001 | LOGIX SMART™ Coronavirus Disease 2019 (COVID-19) Kit | 100T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | CoDx Box (BMS, Bio Molecular Systems) | Co-Diagnostics, Inc | | 12 months | see IFU | nasopharyngeal swabs | For consumables and details of componants refer to IFU | US FDA EUA TGA |
| | DirectDetect™ SARS-CoV-2 Detection Kit | | | | Coyote Bioscience Co Ltd (China) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | Aridia COVID-19 Real-Time PCR Test | | | | CTK Biotech Inc (United States Of America) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| C1020 | Cue COVID-19 Test | | Cue Health Monitoring System | | Cue Health Inc | N Gene | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA replaced by DeNovo approval Health Canada/Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|-------------------------------|---|---|--|----------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 99-57003 | HDPCR™ SARS-CoV-2 Assay | 48oT/kit | Thermo Scientific™ KingFisher™ Flex | 7500 Fast Real-Time PCR System (Applied Biosystems) | ChromaCode, Inc. | N gene (N1 and N2 regions) | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA |
| | | | | QuantStudio7 (Fast Block) instrument (Applied Biosystems) | | | | | | | |
| | | | Roche MagNA Pure-24 | QuantStudio 12k Flex (96-well Fast Block) instrument (Applied Biosystems) | | | | | | | |
| | | | | Bio Molecular Systems Mic qPCR (IDEXX Laboratories) | | | | | | | |
| DA0930 DA0931 DA0932 | Detection Kit for 2019 Novel Coronavirus (2019-nCoV) RNA, (PCR- Fluorescence Probing) | 24T/kit 48T/kit 96T/kit | YHXB No. 20170583, YHXB No. 20150302 (DAAN) | Applied Biosystems™ 7500 Dx Real-Time PCR Instrument | Da An Gene Co., Ltd. of Sun Yat-sen University | ORF1ab and N genes | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL TGA |
| | | | QIAamp Viral RNA Mini Kit, 52906 | Roche LightCycler480 II | | | | | | | |
| | QuickNavi-Flu+COVID19 Ag | | | | Denka Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | Detect Covid-19 Test | 1T/kit | Detect Hub (Model 21101) | | Detecta Inc | ORF1ab | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| MOL4150 | MobileDetect-BIO BCC19 Test Kit | 24T/kit | MD-Bio BCC19 Heater | | DetectaChem LLC | N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|--|--|--------------------|-----------------------|---------------------------------|---------------------------------|--|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| DC-11-0007 | QuantiVirus SARS-CoV-2 Test kit | 24T/kit | Thermo Fisher PureLink™ viral RNA/DNA mini kit | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument | DiaCarta, Inc | N, Orf1ab and E genes | see IFU | see IFU | nasal swabs, nasopharyngeal swabs, oropharyngeal swabs, and sputum | For consumables and details of components refer to IFU | US FDA EUA |
| DC-11-0008 | | 48T/kit | | Applied Biosystems™ 7500 Fast Dx Real-Time PCR Instrument | | | | | | | |
| DC-11-0009 | | 480T/kit | | | | | | | | | |
| DC-11-0017 | QuantiVirus SARS-CoV-2 Multiplex Test kit | 24T/kit | Thermo Fisher PureLink™ viral RNA/DNA mini kit | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument | DiaCarta, Inc | Orf1ab genes | 12 months | see IFU | nasal swabs, nasopharyngeal swabs, oropharyngeal swabs, and sputum | For consumables and details of components refer to IFU | US FDA EUA |
| DC-11-0018 | | 48T/kit | MGI MGISP960 High Throughput Automated Sample Preparation System | Applied Biosystems™ 7500 Fast Dx Real-Time PCR Instrument | | | | | | | |
| DC-11-0019 | | 480T/kit | | Bio-Rad CFX 384 Real-Time PCR Instrument | | | | | | | |
| MOL4150 | Simplexa™ COVID-19 Direct | 24T/kit | LIAISON® MDX | | DiaSorin Molecular | ORF1ab and S gene | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | Health Canada/Interim Order TGA WHO EUL |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|--------------------------|----------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | DTPM COVID-19 RT-PCR Test | 96t/kit | QIAamp Viral RNA Kit (Qiagen) | QuantStudio 5 | DTPM, Inc | S and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | QIAamp 96 Viral RNA Kit (Qiagen) | | | | | | | | |
| | | | Indical Biosciences IndiMag Pathogen Kit | | | | | | | | |
| | | | Omega Biotek Mag-Bind Viral DNA/RNA 96 Kit | | | | | | | | |
| | | | ThermoFisher MagMAX™ Viral/Pathogen II (MVP II) Nucleic Acid Isolation Kit with KingFisher Flex | | | | | | | | |
| | DxLab COVID-19 Test | 24T/kit | DxHub device | | DXLab Inc. | | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| LMP401 | Loopamp New Coronavirus 2019 (SARS-CoV-2) Detection Reagent Kit | 48T/kit | see IFU | | Eiken Chemical Co., Ltd. | Replicase 1B region | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| ENZ-GEN215-0096 | AMPIPROBE® SARS-CoV-2 Assay kit | | GENFLEX platform V1.0 | | Enzo Life Sciences, Inc. | N gene (N1 and N2 regions) | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA |
| | | | QIAsymphony® SP (QIAGEN) | QuantStudio® 5 Real-Time PCR System (Applied Biosystems) | | | | | | | |
| | | | Manual | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|-------------------------------------|------------------|---|---|---|-------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| MP 2606-0125 | EURORealTime SARS-CoV-2 | 25T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | LightCycler® 480 II (Roche) | EUROIMMUN Medizinische Labordiagnostika AG (Germany) | ORF1ab and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order US FDA EUA TGA |
| MP 2606-0225 | | 50T/kit | Prepito Viral DNA-RNA200 Kit (Chemagen) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | | | | | | | |
| MP 2606-0425 | | 100T/kit | Chemagic Viral DNA/RNA 300 Kit H96 | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| MP 2606-0100 | | 100T/kit | | qTOWER3 (Analytik Jena) | | | | | | | |
| MP 2606-0200 | | 200T/kit | | | | | | | | | |
| MP 2606-1000 | | 1000T/kit | | | | | | | | | |
| 11416302 (FTD-114-96) | FTD SARS-CoV-2 | 96T/kit | NucliSENS® easyMAG® System (bioMérieux) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | Fast Track Diagnostics Luxembourg S.á.r.l. (a Siemens Healthineers Company) | ORF1ab and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA PMDA |
| | | | VERSANT® kPCR Molecular System (Siemens Healthineers) | | | | | | | | |
| 11416300 (FTD-114-32) | FTD SARS-CoV-2 | 32T/kit | NucliSENS® easyMAG® System (bioMérieux) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | Fast Track Diagnostics Luxembourg S.á.r.l. (a Siemens Healthineers Company) | ORF1ab and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| 11416284 (FTD-114-96) | | 96T/kit | | | | | | | | | |
| PCSYHF02-a (48 tests/kit); PCSYHF03-a (96 tests/kit) | Fosun COVID-19 RT-PCR Detection Kit | | see IFU | | Fosun Pharma USA Inc. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--------------------------------------|---------------------|--|---|---|-------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | µTASWako SARS-CoV-2 | | see IFU | | FUJIFILM Wako Pure Chemical Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| CV002 | GenePro SARS-CoV-2 Test | | QIAamp DSP Viral RNA Mini Kit (Qiagen) | QuantStudio™ DX (Applied Biosystems) | Gencurix, Inc. | N-gene and E-gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | QIAamp Viral RNA Mini Kit (Qiagen) | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit automated on KingFisher™ Flex Purification System (KingFisher) | | | | | | | | |
| | NeoPlex COVID-19 Detection Kit | 96T/kit | QIAamp DSP Viral RNA Mini Kit (Qiagen) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | GeneMatrix, Inc. | RdRp and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 4PCO052E | GB SARS-CoV-2 Real Time RT-PCR | 100T/kit | QIAamp DSP Viral RNA Mini Kit (Qiagen) | QIAGEN Rotor-Gene® Q real-time PCR cyclcr | General Biologicals Corp | ORF1ab and E gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | EasyScreen™ SARS-CoV-2 Detection Kit | | | | Genetic Signatures Ltd (Australia) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| RPQ021 RPQ022 | Genetron SARS-CoV-2 RNA Test | 50T/kit 100T/kit | QIAamp DSP Viral RNA Mini Kit (Qiagen) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | Genetron Health (Beijing) Co., Ltd. | ORF1ab and N gene | 6 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| EA008212 | ePlex® SARS-CoV-2 Test | 12T/kit | GenMark ePlex instrument and Software | | GenMark Diagnostics, Inc | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------------------------|---|---|----------------------------------|------------------------------------|---------------------------------|---------------------------------|---|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | GS COVID-19 RT-PCR KIT | 96T/kit | QIAamp DSP Viral RNA Mini Kit | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | GenoSensor LLC | ORF1ab, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | 384T/kit | | | | | | | | | |
| CV0202 | COVID-19 RT-Digital PCR Detection Kit | 48T/kit | QIAamp® DSP Viral RNA Mini Kit (Qiagen) | QuantStudio™ 3D Digital PCR System (Applied Biosystems) | Gnomegen LLC | | see IFU | see IFU | nasal, nasopharyngeal, and oropharyngeal swab | For consumables and details of componants refer to IFU | US FDA EUA |
| | Procleix SARS-CoV-2 Assay | 250T/kit 1000T/kit 5000T/kit | Procleix Panther System with Procleix Reagent Preparation Incubator 250 | | Grifols Diagnostic Solutions Inc | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| NAT-01 | RT-PCR IVD MEDICAL DEVICE FOR 2019 NOVEL CORONAVIRUS (SARS-COV-2) NUCLEIC ACID DETECTION | see IFU | see IFU | | HA TECH PTY LTD | ORF1ab Region 1 ORF1ab Region 2 | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| PRD-06419 | Aptima SARS-CoV-2 assay | 250T/kit | Panther System | | Hologic Inc | ORF1ab Region 1 ORF1ab Region 2 | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA TGA PMDA |
| | Panther Fusion SARS-CoV-2 Kit | 96T/kit | Panther/Panther Fusion System | | Hologic Inc | ORF1ab Region 1 ORF1ab Region 2 | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada Interim Order TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--------------------------------------|------------------|--|--|---------------------------|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 351251 | Hymon™ SARS-CoV-2 Test Kit | 96T/kit | | 7500 Dx Real-Time PCR Instrument (Applied Biosystems™) | HymonBio Co. LTD | N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | QuantStudio 5 RT PCR System (Applied Biosystems™) | | | | | | | |
| COV2-E | Smart Detect™ SARS-CoV-2 rRT-PCR Kit | 48T/kit | QIAamp® DSP Viral RNA Mini Kit (Qiagen) | 7500 Fast Dx Real-Time PCR instrument (Applied Biosystems) | InBios International, Inc | ORF1b, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | MagMAX Viral/Pathogen II Nucleic Acid Isolation Kit Hamilton MagEx Star automated liquid handling system | CFX384 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | COV-19 IDx assay | 96T/kit | KingFisher Flex nucleic acid extraction systems QS12 instrument | Applied Biosystems QuantStudio12 Flex (QS12) instrument | Ipsum Diagnostics, LLC | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|-------------------------------|--|--|--|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| JC10223-1NW-50T JC10223-1NW-25T | COVID-19 Coronavirus Real Time PCR Kit | 50T/kit 25T/kit | Viral nucleic acid isolation kit (Bioperfectus Technologies) | Applied Biosystems QuantStudio5 instrument | Jiangsu Bioperfectus Technologies Co Ltd (China) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA TGA WHO EUL |
| | | | QIAamp® Viral RNA Mini Kit (Qiagen) | 7500 Real-Time PCR instrument (Applied Biosystems) | | | | | | | |
| | Novel Coronavirus (SARS-CoV-2) Fast Nucleic Acid Detection Kit | 24T/kit 48T/kit 96T/kit | see IFU | 7500 Real-Time PCR instrument (Applied Biosystems) | Jiangsu CoWin Biotech Co., Ltd. (China) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | KANEKA Direct RT-PCR kit SARS-CoV-2 | | | | KANEKA CORPORATION | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| RV008 | RADI COVID-19 Detection Kit | 100T/kit | QIAamp® DSP Viral RNA Mini Kit (Qiagen) | CFX96 Real-Time PCR Detection System (Bio-rad) | KH Medical Co. Ltd | S gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | SmartAmp SARS-CoV-2 | | | | K.K. DNAFORM | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| KF2019CoV01 | KimForest SARS-CoV-2 Detection Kit v1 | 96T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | StepOne/StepOnePlus Real-Time PCR Systems (Applied Biosystems) | KimForest Enterprise Co., Ltd. | RdRp genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|---|--|---|---------------------------|---------------------------------|---------------------------------|--|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| R6900TD | PowerChek™ 2019-nCoV Real-time PCR Kit | | QIAamp® DSP Viral RNA Mini Kit (QIAGEN) | CFX96 Real-Time PCR Detection System (Bio-Rad) | Kogene Biotech Co Ltd (Korea - Republic of) | RdRp and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA TGA |
| | | | | 7500 Real-Time PCR instrument (Applied Biosystems) | | | | | | | |
| | | | | 7500 Fast Real-Time PCR instrument (Applied Biosystems) | | | | | | | |
| | LabGun™ COVID-19 RT-PCR Kit | 96T/kit | QIAamp® DSP Viral RNA Mini Kit (Qiagen) | 7500 Fast Dx Real-Time PCR instrument (Applied Biosystems) | LabGenomics | RdRp and N genes | see IFU | see IFU | nasopharyngeal swab, anterior nasal swab and midturbinate nasal swab | For consumables and details of componants refer to IFU | US FDA EUA TGA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | TaqPath COVID-19 Combo Kit | | | | Life Technologies Corporation (USA) (see also ThermoFisher) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA PMDA |
| 50-10047 | ARIES® SARS-CoV-2 Assay Kit | 24T/kit | Luminex® ARIES® System | | Luminex Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order |
| I054C0463 | NxTAG® CoV Extended Panel Assay | 96T/kit | bioMérieux® NucliSENS® easyMAG® System | Luminex® MAGPIX® instrument including xPONENT | Luminex Molecular Diagnostics, Inc. | ORF1ab, N Gene and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order |
| | | | bioMérieux® EMAG® System | | | | | | | | |
| GCRNA-COVID-96R | Genecount Covid-19 Rt-Qpcr Assay Kit | | | | Luminultra Technologies Ltd. (Canada) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|------------------------------|------------------|--|---|-----------------|----------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| Lo18180030096 | LumiraDx SARS-CoV-2 RNA STAR | | Qiagen DSP Virus/Pathogen Kit on QIAasymphony SP (Qiagen) | Roche LightCycler 480 II | LumiraDx UK Ltd | ORF1a Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMax Viral/Pathogen Nucleic Acid Isolation Kit (Applied Biosystem) | Applied Biosystems 7500 Fast Dx | | | | | | | |
| | | | QIAamp Viral RNA Mini Kit (Qiagen) | Applied Biosystems QuantStudio 5 | | | | | | | |
| | | | | Agilent AriaMx RT-PCR Instruments | | | | | | | |
| | | | | Agilent Stratagene Mx3005P RT-PCR Instruments | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | | | | | | | | | | | | | | | | |
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| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others | | | | | | | | | | | | | | | |
| L018180130096 | LumiraDx SARS-CoV-2 RNA STAR Complete | | n/a | Applied Biosystems 7500 Fast Dx | LumiraDx UK Ltd | ORF1a Gene | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA | | | | | | | | | | | | | | | |
| | | | | Applied Biosystems QuantStudio 5 | | | | | | | | | | | | | | | | | | | | | | |
| | | | | Applied Biosystems QuantStudio 7 Flex | | | | | | | | | | | | | | | | | | | | | | |
| | | | | Roche LightCycler 480 II | | | | | | | | | | | | | | | | | | | | | | |
| | | | | Bio-Rad CFX96 Touch Real-Time PCR Detection System | | | | | | | | | | | | | | | | | | | | | | |
| | | | | Agilent AriaMx RT-PCR Instruments | | | | | | | | | | | | | | | | | | | | | | |
| | | | | Agilent Stratagene Mx3005P RT-PCR Instruments | | | | | | | | | | | | | | | | | | | | | | |
| BUSGN7101109 | SARS-CoV-2 Fluorescent PCR Kit | 32T/kit | Nucleic Acid Extraction Kit, Manual Version or Nucleic Acid Extraction Kit, Fast Version | 7500 Real-Time PCR Systems with v2.3 software (Applied Biosystems) | Maccura Biotechnology (USA) LLC | ORF1ab, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA | | | | | | | | | | | | | | | |
| BUSGN7102109 | | 64T/kit | | | | | | | | | | | | | | | | | | | | | | | | |
| BUSGN7103109 | | 96T/kit | QIAGEN QIAamp Viral RNA Mini Kit | | | | | | | | | ST-CV19-2SF | MatMaCorp COVID-19 2SF Test | | MatMaCorp Solas 8 Instrument | | Materials and Machines Corporation of America (DBA MatmaCorp, Inc.) | RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA | | MEBRIGHT SARS-CoV-2 Kit | |
| ST-CV19-2SF | MatMaCorp COVID-19 2SF Test | | MatMaCorp Solas 8 Instrument | | Materials and Machines Corporation of America (DBA MatmaCorp, Inc.) | RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA | | | | | | | | | | | | | | | |
| | MEBRIGHT SARS-CoV-2 Kit | | | | Medical & Biological Laboratories Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA | | | | | | | | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
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| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 410700 | Revogene SARS-CoV-2 assay | | REVOGENE SYSTEM | | Meridian | N gene | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA |
| COV4100 | Accula SARS-Cov-2 Test | | Accula™ Dock or the Silaris™ Dock | | Mesa Biotech Inc. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 7K105 7K111 | Veri-Q PCR 316 COVID-19 Detection Kit | 50 test/kit 100 test/kit | Veri-Q System | | MiCoBioMed Co Ltd | ORF3a and N gene target | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA WHO EUL |
| SCFo030 | MicroGEM Sal6830 SARS-CoV-2 Saliva Test | 30T/kit | MicroGEM Sal6830 Point of Care PCR System (SCFMA) | | MicroGEM U.S., Inc. | E and N gene target | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order |
| PN-0205 | DASH SARS-CoV-2/S Test | 768T/kit | DASH Analyzer | | Minute Molecular Diagnostics, Inc. | N Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Smart Gene SARS-CoV-2 | | | | Mizuhomedy Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | MobileDetect Bio BCC19 (MD-Bio BCC19) Test Kit | 12T/kit | MD-Bio heater | | MobileDetect Bio Inc. | E and N gene target | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 300800 | NeuMoDx™ SARS-CoV-2 Test Strip | 96T/kit | NeuMoDx™ 288 Molecular System [500100] or NeuMoDx™ 96 Molecular System [500200] | | NeuMoDx Molecular, Inc. | Nsp2 target and N gene target | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA TGA |
| XC25073 | Sars-Cov-2 Virus Detection Diagnostic Kit | 50T/kit | TAN Bead® extract system (Taiwan Advanced Nanotech) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | Ningbo Health Gene Technologies Co., Ltd (China) | ORF1ab, N and S genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | RNeasy Mini Kit (Qiagen) | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
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| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| DXTM67120 (500RXNS) | 2019-NCoV Taqman Rt-PCR Kit Dx | | | | Norgen Biotek Corp. (Canada) | RdRp and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| DXTM67100 (50RXNS) | 2019-NCoV Taqman Rt-PCR Kit Dx | | | | Norgen Biotek Corp. (Canada) | RdRp and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| DXTM67200 | COVID-19 Taqman Rt-PCR Kit (e/rdrp Genes) Dx | | | | Norgen Biotek Corp. (Canada) | RdRp and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| KT1010001 | Hi-Sense COVID-19 Molecular Testing Kit 1.0 | 96t/kit | TBD | | OnsiteGene, Inc. | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| RDM101-X | Kaira 2019-nCoV Detection Kit | 100T/kit | QIAsymphony DSP Virus/Pathogen Kit on QIAsymphony SP (Qiagen) | 7500 Fast Real-Time PCR System (Applied Biosystems) | OPTOLANE Technologies, Inc. | RdRp and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | QuantStudio5 Flex (QS5) instrument (Applied Biosystems) | | | | | | | |
| | | | | CFX96 Real-Time PCR Instrument (Biorad) | | | | | | | |
| IFMR-45 | GeneFinder™ COVID-19 Plus RealAmp Kit | 100T/kit | QIAamp viral RNA Mini Kit (Qiagen) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | OSANG Healthcare Co., Ltd | RdRp gene, N Gene and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | DNA and Viral NA Small Volume Kit (Roche MagNA Pure 96) | CFX96 Real-Time PCR Instrument (Biorad) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|-----------------------------------|------------------|---|---|----------------------------|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 99-57003 99-57004 | OPTI SARS-CoV-2 RT PCR Test | | Duo instrument (Thermo Scientific) | 7500 Fast Real-Time PCR System (Applied Biosystems) | OPTI Medical Systems, Inc. | N gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | QuantStudio5 Flex (QS5) instrument (Applied Biosystems) | | | | | | | |
| | | | | LightCycler 480 (System II) (Roche) | | | | | | | |
| | | | Flex instrument (Thermo Scientific) | Agilent Mx3005P™ (Agilent) | | | | | | | |
| | | | | Bio Molecular Systems Mic qPCR (IDEXX Laboratories) | | | | | | | |
| | P23 Labs TaqPath SARS-CoV-2 Assay | see IFU | see ThermoFisher TaqPath COVID-19 Combo Kit | | P23 Labs, LLC | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | DetectX-Rv | see IFU | | MiniAmp A37834 (ThermoFisher) | PathogenDx, Inc. | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

SARS-CoV-2 Nucleic Acid Amplification Technologies

| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
|---------------------------------------|---|------------------|--|--|---|------------------------|---------------------------------|---------------------------------|---------------|--|--|
| 2019-nCoV-PCR-AUS | PerkinElmer® New Coronavirus Nucleic Acid Detection Kit | 48T/kit | PerkinElmer® Nucleic Acid Extraction Kits (KN0212) and PreNAT II (SY61)(software version 1.00.06). | Applied Biosystems 7500 Real-Time PCR System | PerkinElmer, Inc. / Suzhou Sym-Bio Lifescience Co Ltd | ORF1ab gene and N gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order WHO EUL |
| | | | chemagic™ Viral DNA/RNA 300 Kit special H96 (CMG-1033, CMG-1033-S) and chemagic™ 360 (2024-0020) with chemagic™ Rod Head Set 96 (CMG371) | Applied Biosystems™ 7500 Fast Dx Real-Time PCR System | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 3 Real-Time PCR System | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 5 Real-Time PCR System | | | | | | | |
| | | | | Analytik Jena qTower3 / qTower3G Real-Time PCR System | | | | | | | |
| | | | | Analytik Jena qTower3 84 / qTower3 84 G Real-Time PCR System | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|--|--|------------------------------------|-------------------------|---------------------------------|---------------------------------|---------------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 82303-U | IntelliPlex SARS-CoV-2 Detection Kit | 96T/kit | QIAmp Viral RNA Mini Kit (Qiagen) | Thermo Fisher MiniAmp Thermal cycler with IntelliPlex™ 1000 πCode Processor (PlexBio; Cat. No. 80033) and PlexBio 100 Fluorescent Analyzer (PlexBio; Cat. No. 80000) | PlexBio Co., Ltd. | RdRp gene, N and E Gene | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 02.01.1019 | FastPlex Triplex SARS-CoV-2 detection kit | 24T/kit | QIAmp Viral RNA Mini Kit (Qiagen) | DropX-2000 Digital PCR System | PreciGenome LLC | RdRp gene and N Gene | see IFU | see IFU | nasopharyngeal swab | For consumables and details of componants refer to IFU | US FDA EUA |
| | Triplelock SARS-Cov-2 Test Strips | | | | Precision Biomonitoring Inc | | see IFU | see IFU | nasopharyngeal swab | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| | ELITe MGB SARS-CoV-2 PCR Detection Kit | | | | Precision System Science Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | LeaDEA VIASURE SARS-CoV-2 PCR Kit | | | | Precision System Science Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| Z-PATH-COVID-19-CE | COVID-19 genesig® Real-Time PCR assay | 96T/kit | GXT DNA/RNA Extraction kit (GenoXtract®, Bruker-HAIN Lifescience GmbH) | Applied Biosystems® 7500 Real-Time PCR System | Primerdesign Ltd | Orf1 ab gene | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL TGA |
| | | | | Bio-Rad CFX Connect™ Real-Time PCR Detection System | | | | | | | |
| | | | | Roche® LightCycler 480 II | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------------------|------------------|--|--|---|----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| Z-COVID-19 (US ONLY) | COVID-19 genesig® Real-Time PCR assay | 96T/kit | GXT DNA/RNA Extraction kit (GenoXtract®, Bruker-HAIN Lifescience GmbH) | Applied Biosystems® 7500 Real-Time PCR System | Primerdesign Ltd | Orf1 ab gene | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Bio-Rad CFX Connect™ Real-Time PCR Detection System | | | | | | | |
| | | | QIAamp Viral RNA Mini kit | Roche® LightCycler 480 II | | | | | | | |
| PCCSKU15261 | PhoenixDx® 2019-nCoV | 50T/kit | RTA Viral RNA Extraction Kit as extraction Kit (RTA Laboratories) | 7500 Fast Real-Time PCR System (Applied Biosystems) | Procomcure Biotech GmbH (Trax Management Services Inc.) | E gene and RdRp gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | QIAamp MinElute Virus Spin Kit (Qiagen) | Rotor-Gene Q (Qiagen) | | | | | | | |
| | | | High Pure Viral RNA Kit (Roche) | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------------|------------------|---|---|---|--------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| PCCSKU15262 | PhoenixDx® SARS-CoV-2 Multiplex | 50T/kit | RTA Viral RNA Extraction Kit as extraction Kit (RTA Laboratories) | 7500 Fast Real-Time PCR System (Applied Biosystems) | Procomcure Biotech GmbH (Trax Management Services Inc.) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMax Viral/Pathogen Nucleic Acid Isolation Kit (Thermo Fisher) | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | QIAamp MinElute Virus Spin Kit (Qiagen) | Rotor-Gene Q (Qiagen) | | | | | | | |
| | | | High Pure Viral RNA Kit (Roche) | DTPPrime5 (DNA Technologie) | | | | | | | |
| | | | SphaeraMag DNA/RNA Isolation Kit (Procomcure) | qTower3G (Analytik Jena) | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 1 Real-Time PCR Instrument (ThermoFisher) | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 3 Real-Time PCR Instrument (ThermoFisher) | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument (ThermoFisher) | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 7 Real-Time PCR Instrument (ThermoFisher) | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|--|------------------------------------|---------------------------------|---------------------------------|----------------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 691223 | QIAstat-Dx Respiratory SARS-CoV-2 Panel | 6 Tests | QIAstat Dx Analyzer System | | QIAGEN GmbH | Orf1b poly gene (Rdrp) and E genes | see IFU | see IFU | nasopharyngeal swab | For consumables and details of components refer to IFU | US FDA EUA Health Canada/Interim Order TGA |
| #1110 with #1105 #1154 with #1155 | Clarifi COVID-19 Test Kit | 480T/kit | Quick-RNA Viral 96 Kit | CFX96 Touch Real-Time Detection System (Biorad) | Quadrant Biosciences Inc. | RdRp genes | see IFU | see IFU | saliva swab specimen | For consumables and details of components refer to IFU | US FDA EUA |
| | | | | CFX384 Touch Real-Time Detection System (Biorad) | | | | | | | |
| | | | | QuantStudio 5 instrument (Applied Biosystems) | | | | | | | |
| 39433 | Quest SARS-CoV-2 rRT-PCR Kit | 96T/kit | Roche MagNA Pure-96 (MP96) | Applied Biosystems 7500 Real Time PCR System | Quest Diagnostics Infectious Disease Inc | Gene N1 & N3 | 12 months | see IFU | see IFU | For consumables and details of components refer to IFU | US FDA EUA |
| | | | Mag-Bind Viral RNA Xpress Kit (Omega Bio-Tek, Cat. M6219-2304) with the MagEx STAR (Hamilton) | | | | | | | | |
| | Quest Diagnostics RC SARS-CoV-2 rRT-PCR Kit | 96T/kit | cobas 6800/8800 (Roche) | | Quest Diagnostics Infectious Disease Inc | ORF1 a/b | 12 months | see IFU | see IFU | Quest Diagnostics Self-Collection Kit for COVID-19 For consumables and details of components refer to IFU | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---------------------------------|--|--|------------------------------------|---------------------------------|---------------------------------|--|---|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | Quest Diagnostics HA SARS-CoV-2 rRT-PCR Kit | 96T/kit | Aptima (Hologic) | | Quest Diagnostics Infectious Disease Inc | ORF1ab Region 1 ORF1ab Region 2 | 12 months | see IFU | nasopharyngeal swabs, oropharyngeal swabs, sputum, BAL, and tracheal aspirates | Quest Diagnostics Self-Collection Kit for COVID-19 For consumables and details of componants refer to IFU | US FDA EUA |
| | Quest Diagnostics PF SARS-CoV-2 rRT-PCR Kit | 96T/kit | Panther Fusion (Hologic) | | Quest Diagnostics Infectious Disease Inc | ORF1ab Region 1 ORF1ab Region 2 | 12 months | see IFU | nasopharyngeal swabs, oropharyngeal swabs, sputum, BAL, and tracheal aspirates | Quest Diagnostics Self-Collection Kit for COVID-19 For consumables and details of componants refer to IFU | US FDA EUA |
| SKU # CE-M120 | Lyra SARS-CoV-2 rRT-PCR Kit | 96T/kit | bioMérieux NucliSENS easyMAG | Applied Biosystems 7500 Real Time PCR System | Quidel Corp. | Orf1ab | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada Interim Order |
| | | | | Applied Biosystems 7500 Fast Dx Real-Time PCR System | | | | | | | |
| | | | | Roche LightCycler 480 | | | | | | | |
| | | | | Qiagen Rotor-Gene Q | | | | | | | |
| | | | | Bio-Rad CFX96 Touch | | | | | | | |
| | | | | Thermofisher QuantStudio 7 Pro | | | | | | | |
| M313 | Solana SARS-CoV-2 Assay | 96T/kit | Solana Instrument | | Quidel Corp. | Orf1ab | see IFU | see IFU | Nasopharyngeal or oropharyngeal specimens | For consumables and details of componants refer to IFU | US FDA EUA Health Canada Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|----------------------|--|---------------------------------------|------------------------|---------------------|---------------------------------|---------------------------------|---------------------------------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| KCCOV19-24 | Rheonix COVID-19 MDx Assay | 96T/kit | Rheonix Encompass MDx® Workstation | | Rheonix | ORF1 a/b | see IFU | see IFU | nasopharyngeal and oropharyngeal swab | For consumables and details of components refer to IFU | US FDA EUA |
| 09175431190 09343733190 | Cobas SARS-CoV-2 RT-PCR Kit | 192T/kit 480T/kit | cobas 6800/8800 | | Roche Diagnostics | ORF1 a/b and E gene | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | US FDA EUA Health Canada/Interim Order PMDA WHO EUL |
| 09500111190 | Cobas SARS-CoV-2 Duo | 192T/kit | cobas 6800/8800 | | Roche Diagnostics | ORF1 a and ORF1 a/b | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | US FDA EUA |
| 09446109190 09448870190 | Cobas SARS-CoV-2 RT-PCR Kit | 192T/kit 480T/kit | cobas 5800/6800/8800 | | Roche Diagnostics | ORF1 a/b and E gene | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | PMDA |
| | cobas Liat SARS-CoV-2 | 20T/kit | cobas Liat System | | Roche Diagnostics | | see IFU | see IFU | see IFU | For consumables and details of components refer to IFU | PMDA |
| 09408592190 | cobas SARS-CoV-2 | 20T/kit | cobas Liat System | | Roche Diagnostics | ORF1ab and N genes | 24 months | see IFU | see IFU | For consumables and details of components refer to IFU | Health Canada/Interim Order PMDA |
| 09408592190 | cobas SARS-CoV-2 | 20T/kit | cobas Liat System | | Roche Diagnostics | ORF1ab and N genes | 18 months | see IFU | see IFU | For consumables and details of components refer to IFU | US FDA EUA |
| S2104F | Novel Coronavirus (2019-nCoV) Nucleic Acid Diagnostic | 24T/kit | QIAamp Virus RNA Mini Kit (cat. # 52904) | Applied Biosystems 7500 Real-Time PCR | Sensaura Bio Tech Inc. | ORF1ab and N | see IFU | see IFU | see IFU | For consumables and details of | US FDA EUA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--------------------------------|------------------|---------------------------------|------------------------------------|--------------|----------------|---------------------------------|---------------------------------|---------------|-------------------------|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 531042 | Kit (PCR-Fluorescence Probing) | 48T/kit | without Extraction | 7500 Real-Time PCR System | BioMérieux | genes | see IFU | see IFU | see IFU | componants refer to IFU | TGA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|--|---------------------|-------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| M-NCOV-01 | STANDARD M nCoV Real-Time Detection kit | 96T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | SD Biosensor | ORF1ab, E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | | | | | | | |
| | | | | Roche LightCycler 480 Real-Time PCR systems | | | | | | | |
| SS-9930 | U-TOP™ COVID-19 Detection Kit | 96T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | Seasun Biomaterials | ORF1ab and N Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | | | | | | | |
| | | | PANAMAX Viral DNA/RNA Extraction Kit performed on the PANAMAX 48 Nucleic Acid Extraction System | | | | | | | | |
| | | | TOP Viral DNA/RNA Extraction Kit | | | | | | | | |
| SS-9920 | AQ-TOP COVID-19 Rapid Detection Kit | 96T/kit | QIAamp DSP Virus Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | Seasun Biomaterials | ORF1ab and N Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 Real-Time PCR System | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|-------------------|--|---|---------------------|------------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| SS-9940 | AQ-TOP COVID-19 Rapid Detection Kit PLUS | 96T/kit | QIAamp DSP Virus Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | Seasun Biomaterials | ORF1ab and N Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 Real-Time PCR System | | | | | | | |
| | | | PANAMAX Viral DNA/RNA Extraction Kit performed on the PANAMAX 48 Nucleic Acid Extraction System | | | | | | | | |
| | | | TOP Viral DNA/RNA Extraction Kit | | | | | | | | |
| RP10243X / RP10252W | Allplex™ 2019-nCoV Assay kit | 100T/kit 124T/kit | Microlab STARlet IVD (Cat. No. 173000-075, Hamilton Co.) | CFX96 Real-Time PCR Instrument (Biorad) | Seegene Inc | RdRp gene, N Gene and E Gene | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | STARMag 96 X 4 Universal Cartridge Kit (Cat. No. 744300.4.UC384, Seegene Inc.) using Microlab NIMBUS IVD instrument (Microlab) | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | | | QIAamp DSP Viral Mini Kit using QIAcube instrument (QIAgen) | Applied Biosystems 7500 & 7500 Fast Dx Real-Time PCR System | | | | | | | |
| | | | Ribospin vRD Viral RNA/DNA Extraction Kit (GeneAll) (manual) | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|---|---|--|-----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | | | MagMAX Viral/Pathogen Nucleic Acid Isolation Kit using KingFisher Flex instrument | | | | | | | | |
| | | | MagNA Pure DNA and Viral NA Small Volume Kit using Roche MagNA Pure 96 | | | | | | | | |
| | | | AdvanSure NA EX Kit (extraction kit) using AdvanSure E3 Instrument System (LG Chem) | | | | | | | | |
| see IFU | Allplex SARS-CoV-2 Fast PCR Assay | see IFU | see IFU | see IFU | Seegene Inc | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| see IFU | Allplex RV Master Assay | see IFU | see IFU | see IFU | Seegene Inc | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| PCSYHF02-a | Fosun COVID-19 RT-PCR Detection Kit | 48T/kit | QIAamp DSP Viral RNA Mini Kit (Qiagen) | 7500 Fast Dx Real-Time PCR Instrument (Applied Biosystems™) | Shanghai Fosun Long March Medical Science Co Ltd (China) | ORF1ab, N and E genes | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| PCSYHF03-a | | 96T/kit | | | | | | | | | |
| PCSYHF | Novel Coronavirus (2019-nCoV) RT-PCR Detection Kit (commercial name: Fosun 2019-nCoV qPCR) | | | | Shanghai Fosun Long March Medical Science Co Ltd (China) | ORF1ab, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|--|--|---|--------------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| GZ-D2RM25 | Novel Coronavirus 2019-nCoV Nucleic Acid Detection Kit (Real-time PCR) | 50T/kit | GenAct NE-48 (Shanghai GeneoDx) | 7500 Real-Time PCR Instrument (Applied Biosystems™) | Shanghai GeneoDx Biotechnology Co., Ltd (China) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | QIAamp DSP Viral RNA Mini Kit (Qiagen) | | | | | | | | |
| KH-G-M-574-48 | Diagnostic kit for SARS-CoV-2 Nucleic acid (Real-time PCR) | 48T/kit | nucleic acid extraction product and equipment of Shanghai Kehua bio-engineering Co., Ltd | Applied Biosystems 7500 Real-Time PCR System | Shanghai Kehua bio-engineering Co., Ltd | ORF1ab and N genes and E genes | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | | CFX96 Real-Time PCR Instrument (Biorad) | | | | | | | |
| | | | | Tianlong Gentier 96E | | | | | | | |
| RR-0485-02 | Novel Coronavirus (2019-nCoV) Real Time Multiplex RT-PCR Kit | 25T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | Shanghai ZJ Bio-Tech Co Ltd (China) | ORF1ab and N genes and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| RR-0479-02 | Novel Coronavirus (2019-nCoV) Real Time Multiplex RT-PCR Kit (Detection for 3 genes) | 25T/kit | nucleic acid extraction product and equipment of Shanghai ZJ Bio-Tech | ABI Prism 7500 | Shanghai ZJ Bio-Tech Co Ltd (China) | ORF1ab and N genes and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | | | QIAamp Virus RNA Mini Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | | | | | | | |
| | | | | SLAN | | | | | | | |
| | | | | MIC POC Dx48 | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
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| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | Sherlock CRISPR SARS-CoV-2 Kit | 96T/kit | PureLink™ Viral RNA/DNA Mini Kit (Thermo Fisher Scientific) | Not required (RT-LAMP and CRISPR Technology used) | Sherlock BioSciences, Inc. | ORF1ab and N genes | 12 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Ampdirect 2019-nCoV detection kit | | | | Shimadzu Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| GNT2011-1 | Ezplex SARS-CoV-2 G Kit | 100T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | CFX96 Real-Time PCR Instrument (Biorad) | SML GENETREE Co., Ltd. | RdRP and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems 7500 Real-Time PCR System | | | | | | | |
| | DiaPlexQ Novel Coronavirus (2019-nCoV) Detection Kit | | | | SolGent Co., Ltd (represented by JK Toxpert) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| ASM-00144 | Spartan COVID-19 System Test Cartridge | see IFU | Spartan COVID-19 System | | Spartan Bioscience Inc. (Canada) | see IFU | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |
| | PlexPCR® SARS-CoV-2 | | | | Speedx Pty Ltd (Australia) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | SGNP nCoV/Flu PCR Detection Kit | | | | SUDx-Biotec Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | PerkinElmer® SARS-CoV-2 Real-time RT-PCR Assay | | | | Suzhou Sym-Bio Lifescience Co Ltd (China) (represented by PerkinElmer) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | 2019-nCoV Fluorescence Detection Real-time RT-PCR Kit | | | | Sysmex Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|--|--|--------------------------|--------------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 80-10284 | T2SARS-CoV-2 Panel | 12T/kit | T2Dx® Instrument | | T2 Biosystems, Inc. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Takara SARS-CoV-2 Direct PCR detection kit | | | | Takara Bio Inc. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | TangenDx SARS-CoV-2 Molecular Test | | Tangen GeneSpark instrument | | Tangen Biosciences, Inc. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 68020 | ExProbe™ SARS-CoV-2 Testing Kit | | QIAamp Virus RNA Mini Kit (Qiagen) | Applied Biosystems 7500 Real-Time PCR System | TBG Biotechnology Corp | RdRp gene, N and E genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | EZbead Viral Extraction Kit with automated EZbead System-32 instrument | TBG Q6000 Real-Time PCR System | | | | | | | |
| PGA4102P1 (liquid) / PGA4102P2 (lyophilized form) | SARS-CoV-2 Nucleic acid detection kit based on Real-Time PCR platform | | | | Tellgen Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|-------------------------------------|------------------|--|---|---|-----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| A47813 | TaqPath COVID-19 Combo Kit Advanced | 200T/kit | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | Thermo Fisher Scientific Inc | ORF1ab, S and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMAX™ Viral/Pathogen II Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | | | | | | | |
| A47814 | TaqPath COVID-19 Combo Kit | 1000T/kit | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | ORF1ab, S and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMAX™ Viral/Pathogen II Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | | | | | | | |
| A47817 | Taqpath COVID-19 Rt-PCR Kit | | | | Thermo Fisher Scientific Inc | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|--|------------------|--|--|---|-----------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| | TaqPath New Coronavirus (SARS-CoV-2) Real-Time PCR Detection Kit | | | | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| A48067 | TaqPath COVID-19 CE-IVD RT-PCR Kit | 1000T/kit | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems 7500 & 7500 Fast & Fast Dx Real-Time PCR System | Thermo Fisher Scientific Inc | ORF1ab, S and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | WHO EUL |
| | | | MagMAX™ Viral/Pathogen II Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems™ QuantStudio 5 & 5 Dx Real-Time PCR Instrument | | | | | | | |
| | | | | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | | | | | | | |
| A49918 | TaqPath™ COVID-19 Pooling Kit | 384T/kit | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | ORF1ab, S and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMAX™ Viral/Pathogen II Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|--|---|---|-----------------------|---------------------------------|---------------------------------|---------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| A51333 | TaqPath COVID-19 RNase P Combo Kit 2.0 | 1T/kit | MagMAX™ Viral/Pathogen II Nucleic Acid Isolation Kit (manual) and automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems™ QuantStudio 5 Flex Real-Time PCR Instrument | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | | | | | | | |
| A51606 | TaqPath COVID-19 FAST PCR Combo Kit 2.0 | 1T/kit | n/a | Applied Biosystems™ QuantStudio 5 Flex Real-Time PCR Instrument | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | | | | | | | |
| A49869 | Amplitude™ Solution with TaqPath COVID-19 High Throughput Combo Kit | 20000/kit | Amplitude™ Solution automated on a Tecan™ Fluent™ 1080 Automation Workstation | Applied Biosystems™ QuantStudio 7 Flex Real-Time PCR Instrument | Thermo Fisher Scientific Inc (see also Life Technologies Corporation) | ORF1ab, S and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA Health Canada/Interim Order |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|--|--|---|------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| TD1100 | COVID-19 RT-PCR PNA Kit | 100T/kit | RNeasy Mini kit (Qiagen) | Applied Biosystems 7500 & 7500 Fast Real-Time PCR System | TNS Co., Ltd (Bio TNS) | RdRp and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | CFX96 Touch Real-Time PCR Detection System (Bio-Rad) | | | | | | | |
| | TRCReady SARS-CoV-2 | | TRCReadyR-80 and Chipset for TRCRR detection reagent | | Tosoh Corporation | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| | Gene Cube SARS-CoV-2 | 48T/kit | QIAmp Viral RNA Mini Kit (Qiagen) | | Toyobo Co., Ltd. | N gene | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | PMDA |
| | Gene Cube HQ SARS-CoV-2 | | | | Toyobo Co., Ltd. | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | PMDA |
| UOL001 | UOL COVID-19 Test | | Uh-Oh Labs Point-of-Care Instrument | | Uh-Oh Labs | | 20 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Diagnostic Kit for Novel-Coronavirus (2019-nCoV) RNA (Isothermal Amplification-Real Time Fluorescence Assay | | | | Ustar Biotechnologies (Hangzhou) Co Ltd (China) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | EasyNat Diagnostic Kit for Novel-Coronavirus (2019-nCoV) RNA (Isothermal Amplification-Real Time Fluorescence Assay | | | | Ustar Biotechnologies (Hangzhou) Co Ltd (China) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| 300681 | ViroKey™ SARS-CoV-2 RT-PCR Test | 4x50T/kit | see IFU | Sentosa® SA201 Real-Time PCR Instrument | Vela Operations Singapore Pte Ltd | Orf1a and RdRp | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA TGA |
| | | | | Applied Biosystems® 7500 Fast Dx Real-Time PCR System | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---|------------------|---|---|--|--------------------|---------------------------------|---------------------------------|---------------|--|--|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| 301068 | ViroKey™ SARS-CoV-2 RT-PCR Test v2.0 | 8x48T/kit | see IFU | Sentosa® SA201 Real-Time PCR Instrument | Vela Operations Singapore Pte Ltd | Orf1a and RdRp | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | | Applied Biosystems® 7500 Fast Dx Real-Time PCR System | | | | | | | |
| | SARS-CoV-2 RT-qPCR Reagent Kit | | | | Wallac Oy (Finland) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| | COVID-19 (SARS-CoV-2) Nucleic Acid Test kit | | | | Wuhan EasyDiagnosis Biomedicine Co Ltd (China) | | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | TGA |
| 801301 | SARS-CoV-2 Test Kit (Real-time PCR) | 48T/kit | Virus RNA Extraction Kit (Xiamen Zeesan) | Applied Biosystems™ QuantStudio 3 real-time PCR | Xiamen Zeesan Biotech Co., Ltd. (China) | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | Lab-Aid Virus RNA Extraction Kit on Lab-Aid 824s Nucleic Acid Extraction System | Bio-Rad CFX96 Real-Time System | | | | | | | |

| SARS-CoV-2 Nucleic Acid Amplification Technologies | | | | | | | | | | | |
|--|---------------------------------------|---------------------|--|--|--------------------------------------|--------------------|---------------------------------|---------------------------------|---------------|--|---|
| Manufacturer Product Catalogue number | BinaxNOW COVID-19 Ag Card | Reference detail | Platform (Extraction equipment) | Platform (amplification equipement) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
| SC-COVID19-20 SC-COVID19-100 | COVID-19 Nucleic Acid RT-PCR Test Kit | 20T/kit 100T/kit | QIAamp Virus RNA Mini Kit (Qiagen) | Applied Biosystems® 7500 Fast Real-Time PCR System | ZhuHai Sinochips Bioscience Co., Ltd | ORF1ab and N genes | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | | | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit automated on KingFisher™ Flex Purification System (KingFisher) | Applied Biosystems® 7500 Fast Dx Real-Time PCR System | | | | | | | |
| R3011 R3011-1K R3011-10K | Quick SARS-CoV-2 rRT-PCR Kit | | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit automated on KingFisher™ Flex Purification System (KingFisher) | Bio-Rad CFX96 Touch Real-Time PCR Detection System | Zymo Research Corp | N gene | see IFU | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA Health Canada/Interim Order |
| | | | MagMAX™ Viral/Pathogen Nucleic Acid Isolation Kit manual (KingFisher) | Applied Biosystems™ QuantStudio 5 Real-Time PCR Instrument | | | | | | | |
| N/A- NOT APPLICABLE | | | | | | | | | | | |
| Disclaimer:The Global Fund does not endorse or warrant the fitness of any product on the List for a particular purpose. In addition, the Global Fund assumes no responsibility for any misstatement or omission from the list and directs Principal Recipients of Global Fund grants to conduct their own independent confirmation that the information on a given product on the list is accurate before relying on it to make a purchase order for that product, and to ensure that any purchase is in compliance with all the requirements of the Global Fund’s quality assurance policy. The Global Fund does not warrant or represent that the products listed have obtained regulatory approval for use in any particular country of the world, or that their use is otherwise in accordance with the national laws and regulations of any country, including, but not limited to, intellectual property laws. The Global Fund disclaims any and all liability and responsibility for any injury, death, damage or loss of any kind whatsoever that may arise as a result of, or in connection with the procurement, distribution and use of any product included in the list. | | | | | | | | | | | |

List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)

SARS-CoV-2 Gene Sequencing Technologies (only with the intended use to diagnoses acute infections are included)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Reference detail | Platform (Extraction equipment) | Platform (amplification equipment) | Manufacturer | Detection type | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility criteria WHO EUL or others |
|---------------------------------------|----------------------------|------------------|--|------------------------------------|-----------------|-------------------|---------------------------------|---------------------------------|---------------|--|---|
| | BinaxNOW COVID-19 Ag Card | 192T/kit | Clear Dx™ system comprising Hamilton STAR robotic platform and Oxford Nanopore GridION Sequencer and ALPAQUA Magnum FLX on deck magnet | | Clear Labs, Inc | Sars-Cov-2 Genome | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 20043675 | Illumina® COVIDSeq™ Test | 3072T/kit | NovaSeq 6000 Sequencing System | | Illumina | Sars-Cov-2 Genome | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | Health Canada/Interim Order US FDA EUA PMDA |
| | | | NextSeq 500 Sequencing System | | | | | | | | |
| | | | NextSeq 550 Sequencing System | | | | | | | | |
| | | | NextSeq 550Dx Sequencing System | | | | | | | | |

N/A- NOT APPLICABLE

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List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)

A new technology for COVID-19 detection has become available that is much simpler and faster to perform than currently-recommended nucleic acid amplification tests (NAAT), like PCR. This method relies on direct detection of SARS-CoV-2 viral proteins in nasal swabs and other respiratory secretions using a lateral flow immunoassay (also called an RDT) that gives results in < 30 minutes. Though these antigen detection RDTs (Ag-RDTs) are substantially less sensitive than NAAT, they offer the possibility of rapid, inexpensive and early detection of the most infectious COVID-19 cases in appropriate settings. For more detailed technical advice, please, consult the WHO Interim guidance available at: <https://www.who.int/publications/i/item/antigen-detection-in-the-diagnosis-of-sars-cov-2-infection-using-rapid-immunoassays>
https://www.who.int/publications/i/item/WHO-2019-nCoV-Ag-RDTs-Self_testing-2022.1

SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING
(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|--|--|---|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|--|-------------------------------|
| 41FK501 41FK701 41FK801 41FK901 | BinaxNOW COVID-19 Ag Card | 1Test/kit 4Tests/kit 10Tests/kit 20Tests/kit | n/a | Abbott Rapid Diagnostics Jena GmbH (Germany) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| 41FK51 41FK71 41FK81 41FK91 | Panbio COVID-19 Antigen Self-Test | 1Test/kit 4Tests/kit 10Tests/kit 20Tests/kit | n/a | Abbott Rapid Diagnostics Jena GmbH (Germany) | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | TGA WHO EUL |
| 195-100 195-105 | BinaxNOW COVID-19 Ag Card Home Test | 1Test/kit 2Tests/kit | n/a | Abbott Diagnostics Scarborough, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 22 months | see IFU | nasal | For consumables and details of components refer to IFU | US FDA EUA |
| 195-140 195-160 195-170 | BinaxNOW COVID-19 Antigen Self Test | 1Test/kit 2Tests/kit 10Tests/kit | n/a | Abbott Diagnostics Scarborough, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 22 months | see IFU | nasal | Visual read | US FDA EUA |
| LO31-13025, LO31-13035, LO31-13045, LO31-13055, LO31-13065, LO31-13075, LO31-013085, LO31-13095 | SARS-CoV-2 Antigen Rapid Test (Self-Testing) | | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | WHO EUL |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|--|-------------------------------------|---|---|-----------------------------------|---------------------------------|----------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| Lo31-118M5 Lo31-118P5 Lo131-118R5 | Flowflex SARS-CoV-2 Antigen Rapid Test (Self-Testing) | 1Test/kit 2Tests/kit 5Tests/kit | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | ANSM |
| Lo31-118L5 Lo31-118M5 Lo31-118P5 Lo131-118R5 Lo131-118Z5 Lo131-13015 | Flowflex SARS-CoV-2 Antigen Rapid Test (Self-Testing) | | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| | Flowflex SARS-CoV-2 Antigen Rapid Test (Self-Testing) | | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasopharyngeal | Visual read | TGA |
| Lo31-118B5 Lo31-125M5 Lo31-125N5 Lo31-125P5 | Flowflex SARS-CoV-2 Antigen Rapid Test (Self-Testing) (Alternate brand name: On/Go One COVID-19 Antigen Home Test) | | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | 21 months | see IFU | nasal | Visual read | US FDA EUA |
| Lo31-118B5 Lo31-125M5 Lo131-125N5 Lo131-125P5 | FlowFlex COVID-19 Antigen Home Test | 1Test/kit 2Tests/kit 5Tests/kit 25Tests/kit | n/a | Acon Biotech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | Health Canada/Interim Order |
| Lo31-126H5 Lo31-126J5 Lo31-126K5 Lo31-126L5 | FlowFlex COVID-19 Antigen Home Test | | n/a | Acon Laboratories, Inc. (United States) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| ICO-3000 ICO-3000+ ICO-3000C1 ICO-3000C1+ ICO-3000C40 ICO-3000C40+ ICO-3000C5 ICO-3000C5+ | iHealth COVID-19 Antigen Rapid Test | 2T/kit 5T/kit 40T/kit | n/a | Andon Medical Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 9 months | see IFU | nasal | Visual read | Health Canada/Interim Order |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|---------------------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|--|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Advin COVID-19 Antigen Test @Home | 1T/kit 2T/kit 5T/kit 25T/kit | n/a | Advin Biotech, Inc. 10237 Flanders Ct. San Diego, CA 92121 | SARS-CoV-2 nucleocapsid protein antigen | 15 months | see IFU | nasal | Visual read | USFDA |
| PN-0003KT40 | NIDS® COVID-19 ANTIGEN HOME TEST | 1Tests/kit | n/a | ANP Technologies Inc. | SARS-CoV-2 nucleocapsid protein antigen | 11 months | see IFU | nasal | Visual read | US FDA EUA |
| 1067-2 | OSOM COVID-19 Antigen Rapid Test | 2T/kit | n/a | ANP Technologies Inc. (distributed by SEKISUI Diagnostics LLC) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| 810-00012 Rev B | Metrix COVID-19 Test | 2Tests/kit | n/a | Aptitude Medical Systems | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | US FDA EUA |
| A03-50-422S1 A03-50-422S5 A03-50-422S25 | COVID-19 Antigen Home Test | | n/a | Artron Laboratories Inc. (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | see IFU | Visual read | Health Canada/Interim Order |
| COV-S35Pen | Ecotest COVID-19 Antigen Saliva Test kit (COV-S35Pen) | | n/a | Assure Tech (Hangzhou) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |
| COV-S23010H1 COV-S23010H2 COV-S23010H4 COV-S23010H5 COV-S23010H25 | Fastep COVID-19 Antigen Home Test | | n/a | Azure Biotech Inc. | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | US FDA EUA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|--|---|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|------------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Fastep COVID-19 Antigen Pen Home Test | 1test/kit | n/a | Azure Biotech Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| HGCG134S0101 | Hotgen COVID-19 Antigen Home | 1Tests/kit | n/a | Beijing Hotgen Biotech Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | 6 months | see IFU | nasal | Visual read | US FDA EUA |
| 859261/859262/859263 | SARS-CoV-2 Virus Antigen Detection Kit (Colloidal Gold Method) - Self-test | | n/a | BGI Europe A/S (Denmark) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | Hough COVID-19 Home Test | | n/a | BIOHIT HealthCare (Hefei) Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| 90-1119 90-1121 | Istatis COVID-19 Antigen Home Test | | n/a | Biolytical Laboratories Inc. (Canada) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | Health Canada/Interim Order |
| 859256/859257/859258 | BIOSYNEX - Autotest antigénique COVID-19 Ag | 1Tests/kit | n/a | Biosynex Swiss SA | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | ANSM |
| 859261/859262/859263 | BIOSYNEX - Autotest antigénique COVID-19 Ag | 5Tests/kit | n/a | Biosynex Swiss SA | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | ANSM |
| BTK-H2201, BTK-H2202, BTK-H2205, BTK-H2220 | Bio-Self COVID-19 Antigen Home Test | 1Tests/kit 2Tests/kit 5Tests/kit 20Tests/kit | n/a | BioTeke USA, LLC | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | US FDA EUA |
| COV-19CSHC1 COV-19CSHC2 COV-19CSHC5 COV-19CSHC25 | Rapid Response COVID-19 Antigen Rapid Test Cassette - At Home | | n/a | Btnx Inc. | SARS-CoV-2 nucleocapsid protein antigen | 11 months | see IFU | nasal | Visual read | Health Canada/Interim Order TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|--|---|-------------------------------------|----------------------------|---|-----------------------------------|---------------------------------|--------------------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | CorDx COVID-19 Ag Test | 2 tests/kit | n/a | CorDX, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 15 months | see IFU | see IFU | Visual read | US FDA EUA |
| R0182CHT | ImmuView COVID-19 Antigen Home Test | 2 tests/kit 4 tests/kit 20 tests/kit | n/a | CTK Biotech Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| R0182CST | OnSite COVID-19 Ag Self Test (Nasal Swab), 1Test/Kit | 1 tests/kit | n/a | CTK Biotech Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA WHO EUL |
| R0182CST | OnSite COVID-19 Ag Self Test (Nasal Swab), 2Test/Kit | 2 tests/kit | n/a | CTK Biotech Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA WHO EUL |
| R0182CST | OnSite COVID-19 Ag Self Test (Nasal Swab), 5Test/Kit | 5 tests/kit | n/a | CTK Biotech Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| R0182CST-1T, R0182CST-2T, R0182CST-5T, R0182CST-20T | OnSite COVID-19 Ag Self Test (Nasal Swab) | 1Test/kit 2Test/kit 5Test/kit 20Test/kit | n/a | CTK Biotech Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | WHO EUL |
| I-SRS-C-01 | Ellume COVID-19 Home Test | | n/a | Ellume Limited | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | Mid-turbinate nasal swab | Visual read | US FDA EUA |
| ATG 900-207 ATG 900-208 ATG 900-210 | Covclear COVID-19 Rapid Antigen Test | | n/a | Empowered Diagnostics, Llc | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | Health Canada/Interim Order |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|-------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | FUJI DRI-CHEM IMMUNO AG Handy COVID-19 Ag (general use) | | n/a | Fujifilm Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | PMDA |
| | Genabio COVID-19 Rapid Self-Test Kit | | n/a | Genabio Diagnostics Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| | Covid-19 Ag Home Test | | n/a | GenBody Inc | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| COV-19NS1 COV-19NS2 COV-19NS4 COV-19NS5 | Swiftswab COVID-19 Antigen Self Test | | n/a | Gene Biotechnology Enterprises Ltd | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| | SARS-CoV-2 Antigen Test Kit (Colloidal Gold) | | n/a | Genrui Biotech Inc (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | V-Chek COVID-19 Antigen Saliva Test | | n/a | Guangzhou Decheng Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |
| | WHISTLING TEST 2019-nCoV Saliva Ag EASY TEST | | n/a | Guangzhou Decheng Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |
| W634P0013 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 20T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA WHO EUL |

SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING

(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|---------------------------------------|--|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| W634P0014 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 100T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA WHO EUL |
| W634P0015 | Wondfo 2019-nCoV Antigen Test (Lateral Flow Method) | 20T/kit | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA WHO EUL |
| INCP-502H | JusChek SARS-CoV-2 Antigen Rapid Test (Nasal Swab) INCP-502H Self Test | | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| ICOV-802H | JusChek COVID-19 Antigen Rapid Test (Oral Fluid) ICOV-802H | | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | oral | Visual read | TGA |
| | GSD NovaGen SARS CoV-2 Ag Rapid Test (Nasal swab) Self-testing | | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| INCP-502H01 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H02 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H05 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |

SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING

(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|---------------------------------------|---|-------------------------|-------------------------------------|---------------------------------|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| INCP-502H07 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H10 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H20 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H25 | SARS-CoV-2 Antigen Rapid Test | | | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | 19 months | see IFU | nasal | Visual read | Health Canada |
| INCP-502H | All Test SARS-CoV-2 Antigen Rapid Test (Nasal Swab) Self-Test (INCP-502H) | 1 tests/kit | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| ICOV-802H | My Covid Test Antigen Rapid Test (Oral Fluid) Self-Test (ICOV-802H) | | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | oral | Visual read | TGA |
| ICOV-802H | All Test COVID-19 Antigen Rapid Test (Oral Fluid) Self-Test (ICOV-802H) | | n/a | Hangzhou Alltest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | oral | Visual read | TGA |
| | RightSign COVID-19 Antigen Rapid Test Cassette (Nasal Swab) | | n/a | Hangzhou Biotest Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|--|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Clungene Covid-19 Antigen Rapid Test for self-testing | | n/a | Hangzhou Clongene Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | LYHER Novel Coronavirus (Covid-19) Antigen Test Kit (colloidal Gold) Self-Test | | n/a | Hangzhou Laihe Biotech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Realy Novel Coronavirus (SARS-Cov-2) Antigen Rapid Test Device (nasal swab) | | n/a | Hangzhou Realy Tech Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Medriva COVID-19 Rapid Antigen Self-Test | | n/a | Hangzhou Sejoy Electronics & Instruments Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-CoV-2 Antigen Saliva Lolly Test (without software) | | n/a | Hangzhou Sejoy Electronics & Instruments Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-CoV-2 Antigen Rapid Test Cassette | | n/a | Hangzhou Sejoy Electronics & Instruments Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-CoV-2 Antigen Saliva Lolly Test | | n/a | Hangzhou Testsea Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Cellife Covid-19 Antigen Test Cassette for self-testing | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|--------------------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|---|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | GENEDIAN COVID-19 Antigen Test Cassette (Self Test) | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Nuex Medical COVID-19 Antigen Test Cassette | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Medriva COVID-19 Rapid Antigen Test | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SecurePlus COVID-19 Antigen Test Cassette | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Testsealabs Rapid test kit COVID-19 Antigen Test Cassette | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Wellwisse COVID-19 ANTIGEN TEST CASSETTE | | n/a | Hangzhou Testsea Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-COV-2 RAPID ANTIGEN TEST KIT RAT-01 (SELF-TEST) | | n/a | HA TECH PTY LTD | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| GCCOV-502a-H1US GCCOV-502a-H2US GCCOV-502a-H4US GCCOV-502a-H5US | CLINITEST Rapid COVID-19 Antigen Self-Test | 1T/kit 2T/kit 4T/kit 5T/kit | n/a | Healgen Scientific Limited Liability Company (distributed by Siemens Healthineer) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | US FDA EUA Health Canada/Interim Order |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|---------------------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| ACOVGS-7002 ACOVGS-7005 ACOVGS-7025 | Humasis COVID-19 AG Home Test | | n/a | Humasis Co., Ltd. | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| | Celltrion DiaTrust COVID-19 Ag Home Test | 1Test/kit 2Tests/kit 5Tests/kit | n/a | Humasis Co., Ltd. (distributed by Celltrion USA, Inc) | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | nasal | Visual read | US FDA EUA |
| CT-P60 D-2 05 | Celltrion DiaTrust COVID-19 Ag Home Test | 25Tests/kit | n/a | Humasis Co., Ltd. (distributed by Celltrion USA, Inc) | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | nasal | Visual read | US FDA EUA |
| ICO-3000 ICO-3001 ICO-3002 | iHealth COVID-19 Antigen Rapid Test (Alternate brand name: GoodToKnow COVID-19 Antigen Rapid Test) | 2T/kit 5T/kit 40T/kit | n/a | iHealth Labs, Inc | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | US FDA EUA |
| CFPT-17 | Swab-N-Go Home Test COVID-19 Ag | 2T/kit | n/a | Immunostics Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| CAGS-1 CAGS-2 CAGS-5 CAGS-20 | SCoV-2 Ag Detect Rapid Self-Test | 1T/kit 2T/kit 5T/kit 20T/kit | n/a | InBios International Inc | SARS-CoV-2 nucleocapsid protein antigen | 13 months | see IFU | nasal | Visual read | US FDA EUA |
| | InnoScreen COVID-19 Antigen Rapid Test Device (Self Test) | | n/a | Innovation Scientific Pty Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Gardian T3 COVID-19-SARS-CoV-2 Antigen Saliva Self Test | | n/a | Jiangsu Medomics Medical Technology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |

SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING

(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|---------------------------------------|--|---------------------------------------|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| | Orawell COVID-19 Ag Rapid saliva test device (Self-test) | | n/a | Jiangsu Well Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |
| | SARS-CoV-2 Antigen Rapid Test Kit (Colloidal Gold) | | n/a | JOYSBIO (Tianjin) Biotechnology Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-CoV-2 Antigen Rapid Test Kit (Immunochromatography) | | n/a | Labnovation Technologies Inc (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Antigen Assay Kit | | n/a | Maccura Biotechnology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| 94311 95677 97382 92020 | MaximBio ClearDetect COVID-19 Antigen Home Test | 1T/kit 2T/kit 4T/kit 25T/kit | n/a | Maxim Biomedical, Inc | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | US FDA EUA |
| MS-TRA-SD-1 MS-TRA-SD-5 | Medsup COVID-19 Antigen Rapid Test (self-testing) | | n/a | Medsup Medical (Canada) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | see IFU | Visual read | Health Canada/Interim Order |
| | COVIFIND COVID-19 Antigen Self-Test | | n/a | Meril Diagnostics Pvt Ltd (India) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | COVI-Go SARS-CoV-2 Ag Self-Test | | n/a | Mologic, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|--|-------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|---|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | Rapid SARS-COV-2 Antigen Test Card Self-test (with pre - filled buffer) | | n/a | MP Biomedicals Asia Pacific Pte Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | Rapid SARS-COV-2 Antigen Test Card Self-test | | n/a | MP Biomedicals Asia Pacific Pte Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| MD-8150 | Nano-Check COVID-19 Antigen At-Home Test | 1T/kit | n/a | Nano-Ditech Corporation | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| | Synthgene COVID-19 Antigen Rapid Test Kit | | n/a | Nanjing Synthgene Medical Technology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | SARS-COV-2 Nucleocapsid (N) Antigen Rapid Detection Kit (Colloidal gold method) For Nasal Swab Self-Test | | n/a | Nanjing Synthgene Medical Technology Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |
| | VTRUST SARS-CoV-2 Ag | | n/a | NIPRO CORPORATION | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | PMDA |
| | InteliSwab COVID-19 Rapid Test Rx | | n/a | OraSure Technologies, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | US FDA EUA |
| 1001-0616.001 | InteliSwab COVID-19 Rapid Test | 1T/kit | n/a | OraSure Technologies, Inc. | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | US FDA EUA Health Canada/Interim Order |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|--|-------------------------------------|--|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | OHC COVID-19 Antigen Self Test (Alternate brand name: QuickFinder COVID-19 Antigen Self Test) | 1T/kit 2T/kit 4T/kit 25T/kit | n/a | OSANG Healthcare Co., Ltd. (distributed by OSANG LLC) | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | US FDA EUA |
| COVo4ST-1T COVo4ST-2T COVo4ST-3T COVo4ST-5T COVo4ST-25T COVo4ST-50T | Pcl Self Test - COVID 19 AG | 1T/kit 2T/kit 3T/kit 5T/kit 25T/kit 50T/kit | n/a | Pcl Inc. (South Korea) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | Health Canada/Interim Order |
| P0038 P0039 P0040 P0041 | INDICAID COVID-19 Rapid Antigen At-Home Test | 2T/kit 4T/kit 12T/kit 24T/kit | n/a | PHASE Scientific International, Ltd. (distributed by Precision for Medicine) | SARS-CoV-2 nucleocapsid protein antigen | 15 months | see IFU | nasal | Visual read | US FDA EUA |
| | Gmate COVID-19 Ag Saliva | | n/a | Philosys Co Ltd | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | saliva | Visual read | TGA |
| 33301 | Status COVID-19 Antigen Rapid Test for Home Use | 1T/kit | n/a | Princeton BioMeditech Corp. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| 20431 20402 20427 20398 | QuickVue At-Home OTC COVID-19 Test | 1T/kit 2T/kit 10T/kit 25T/kit | n/a | Quidel Corporation | SARS-CoV-2 nucleocapsid protein antigen | 12 months | see IFU | nasal | Visual read | US FDA EUA |
| 20410 20411 20436 | QuickVue At-Home OTC COVID-19 Test | | n/a | Quidel Corporation | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| | SARS-CoV-2 Antigen Rapid Test | | n/a | Qingdao Hightop Biotech Co Ltd (China) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | see IFU | Visual read | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|---|-----------------------------|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|---------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | BIOCREDIT Covid-19 antigen test kit Nasal (for general use) | | n/a | Rapigen, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | PMDA |
| | Salixium COVID-19 Rapid Antigen Test | | n/a | Reszon Diagnostics International Sdn Bhd Malaysia | | | see IFU | saliva/nasal | | TGA |
| | SARS-CoV-2 Rapid Antigen Test II (for general use) | | n/a | Roche Diagnostics | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | PMDA |
| | Pilot COVID-19 At-Home Test | | n/a | SD Biosensor, Inc. | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | | US FDA EUA |
| 9901-NCOV-06G | SARS-CoV-2 Antigen Self Test Nasal | | n/a | SD Biosensor, Inc | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order TGA |
| 09COV120H 09COV120J | Standard I-q COVID-19 AG Home Test | | n/a | SD Biosensor, Inc | SARS-CoV-2 nucleocapsid protein antigen | 24 months | see IFU | nasal | Visual read | Health Canada/Interim Order |
| 99COV200-EN02 99COV200L-EN02 99COV200D-EN02 (use Roche order nbrs) | COVID-19 At-Home Test | 1T/kit 4T/kit 25T/kit | n/a | SD Biosensor, Inc (distributed by Roche) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | US FDA EUA |
| | SARS-CoV-2 Ag Self test kit | | n/a | Shenzhen Watmind Medical Co- Ltd China | | see further product documentation | see IFU | nasal | | TGA |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|--|---|-------------------------------------|---|---|-----------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| G86256 G86277 G86278 G86279 G86280 G86273 | GLINE-2019-nCoV Ag (Self-Test) | 1 T/kit 5 T/kit 7 T/kit 10 T/kit 15 T/kit 20 T/kit | | Shenzhen YHLO Biotech Co., Ltd | | see further product documentation | see IFU | nasal | | WHO EUL |
| | SGTi-flexCOVID-19 Ag | | n/a | Sugentech Inc Republic of Korea | | see further product documentation | see IFU | nasal | | TGA |
| | SureScreen SARS-CoV-2 Antigen Rapid test Cassette (Nasal Swab) (Gold) | | n/a | SureScreen Diagnostics Ltd (United Kingdom) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | SureScreen Diagnostics SARS-CoV-2 Antigen Rapid Test Cassette (Nasal Swab) Gold) with RhinoSwab Junior | | n/a | SureScreen Diagnostics Ltd (United Kingdom) | SARS-CoV-2 nucleocapsid protein antigen | see further product documentation | see IFU | nasal | Visual read | TGA |
| | Speedy Swab Rapid COVID-19 Antigen Self-Test | | n/a | Watmind USA | SARS-CoV-2 nucleocapsid protein antigen | 21 months | see IFU | nasal | Visual read | US FDA EUA |
| 1N40C5-1-CA 1N40C5-2-CA 1N40C5-4-CA 1N40C5-5-CA 1N40C5-8-CA 1N40C5-10-CA 1N40C5-20-CA 1N40C5-40-CA | Rapid SARS-CoV-2 Antigen Test Card | 1T/kit 2T/kit 4T/kit 5T/kit 8T/kit 10T/kit 20T/kit 40T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 18 months | see IFU | nasal | Visual read | Health Canada/Interim Order |

| SARS-CoV-2 Antigen Rapid Diagnostic Tests for SELF TESTING (Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment) | | | | | | | | | | |
|--|------------------------------------|---|-------------------------------------|--|---|---------------------------------|---------------------------------|---------------|-------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| 1N40C5-1-US 1N40C5-2-US 1N40C5-4-US 1N40C5-5-US 1N40C5-8-US 1N40C5-10-US 1N40C5-20-US 1N40C5-40-US | Rapid SARS-CoV-2 Antigen Test Card | 1T/kit 2T/kit 4T/kit 5T/kit 8T/kit 10T/kit 20T/kit 40T/kit | n/a | Xiamen Boson Biotech Co., Ltd. (China) | SARS-CoV-2 nucleocapsid protein antigen | 6 months | see IFU | nasal | Visual read | US FDA EUA |
| N/A- NOT APPLICABLE | | | | | | | | | | |
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List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)**Important Precautionary statements:**

Based on current data, WHO does not recommend the use of antibody-detecting rapid diagnostic tests for patient care but encourages the continuation of work to establish their usefulness in disease surveillance and epidemiologic research. (<https://www.who.int/news-room/commentaries/detail/advice-on-the-use-of-point-of-care-immunodiagnostic-tests-for-covid-19>)

SARS-CoV-2 Antibody Rapid Diagnostic Tests

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|---------------------------------------|---|-------------------------|-------------------------------------|-----|----------------------------------|---------|---------------------------------|---------------------------------|---------------|--|-------------------------------|
| L031-11711 | BinaxNOW COVID-19 Ag Card | see IFU | n/a | n/a | ACON Laboratories, Inc | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| A-RAPCOV01 | RapCov™ Rapid COVID-19 Test | see IFU | n/a | n/a | Advaite | IgG | see IFU | see IFU | see IFU | | US FDA EUA |
| COV-W23M | Assure COVID-19 IgG/IgM Rapid Test Device | see IFU | n/a | n/a | Assure Tech. (Hangzhou Co., Ltd) | IgG/IgM | see IFU | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |
| | Assure COVID-19 IgG/IgM Rapid Test Device rebranded as Ecotest COVID-19 IgG/IgM Rapid Test Device | see IFU | n/a | n/a | Assure Tech. (Hangzhou Co., Ltd) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | Assure COVID-19 IgG/IgM Rapid Test Device rebranded as Fastep COVID-19 IgG/IgM Rapid Test Device | see IFU | n/a | n/a | Assure Tech. (Hangzhou Co., Ltd) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|---|-------------------------|-------------------------------------|-----|--|---------|---------------------------------|---------------------------------|---------------|----------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | NOVA Test® COVID-19 IgG/IgM Antibody Test (Colloidal Gold) | see IFU | n/a | n/a | Atlaslink Beijing Technology Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| | AtomoRapid™ COVID-19 IgM/IgG Antibody Test | see IFU | n/a | n/a | Atomo Diagnostics Limited | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| WJ-2710, WJ-2750 | Wantai SARS-CoV-2 Ab Rapid Test kit | 10 50 | n/a | n/a | Beijing Wantai Biological Pharmacy Enterprise Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA TGA |
| B251C | Tell Me Fast Novel Coronavirus (COVID-19) IgG/IgM Antibody Test | | n/a | n/a | Biocan Diagnostics Inc. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| RTA0203 | SARS-CoV-2 IgM/IgG Antibody Test Kit | 25 | n/a | n/a | Biohit Healthcare (Hefei) Co., Ltd. | IgG/IgM | see IFU | 2 to 30°C | see IFU | | US FDA EUA TGA |
| 90-1092 90-1098 | Insti COVID-19 Antibody Test | see IFU | n/a | n/a | Biolytical Laboratories Inc. (Canada) | IgG/IgM | 12 | see IFU | see IFU | | Health Canada/Interim Order |
| COV-13C25 | Rapid Response COVID-19 IgG/IgM Rapid Test Device | see IFU | n/a | n/a | Btnx Inc. (Canada) | IgG/IgM | see IFU | see IFU | see IFU | | Health Canada/Interim Order |
| | COVID-19 IgG/IgM Rapid Test | see IFU | n/a | n/a | Btnx Inc. (Canada) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|--|-------------------------|-------------------------------------|-----|--|-------------|---------------------------------|---------------------------------|--|----------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| 5515C025 | qSARS-CoV-2 IgG/IgM Rapid Test | 25 | n/a | n/a | Cellex Inc (United States of America) | IgG/IgM | 12 | 2 to 30°C | serum, plasma (EDTA or citrate), or venipuncture whole blood | | TGA revoked US FDA EUA |
| 5515C050 | | 50 | | | | | | | | | |
| 5515C100 | | 100 | | | | | | | | | |
| R0180C | OnSite COVID-19 IgG/IgM Rapid Test (Aria COVID-19 IgG/IgM Rapid Test) | see IFU | n/a | n/a | CTK Biotech Inc (USA) | IgG/IgM | see IFU | see IFU | see IFU | | TGA revoked US FDA EUA |
| 2039 | CovAb SARS-CoV-2 Ab Test | see IFU | n/a | n/a | Diabetomics, Inc. | IgG/IgA/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| W195 | SARS-CoV-2 Antibody Test (Lateral Flow Method) | see IFU | n/a | n/a | Guangzhou Wondfo Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| INCP-402 | 2019-n-CoV IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Alltest Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA removed from US FDA EUA |
| | COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Biotest Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA removed from US FDA EUA |
| | RightSign™ COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Biotest Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | RightSign™ COVID-19 IgG/IgM Rapid Test Cassette rebranded to CoronaCHEK COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Biotest Biotech Co Ltd (China) distributed by CLIAwaived Inc. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|--|-------------------------|-------------------------------------|-----|---|---------|---------------------------------|---------------------------------|---------------|------------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | RightSign™ COVID-19 IgG/IgM Rapid Test Cassette rebranded to Premier Biotech COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Biotest Biotech Co Ltd (China) distributed by Premier Biotech Inc. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Hangzhou Clongene Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| 303002 | LYHER Novel Coronavirus (2019-nCoV) IgM/IgG Antibody Combo Test Kit (Colloidal Gold) | see IFU | n/a | n/a | Hangzhou Laihe Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | IFU update | US FDA EUA |
| | LYHER Novel Coronavirus (2019-nCoV) IgM/IgG Antibody Combo Test Kit (Colloidal Gold) rebranded to QUICKKIT Novel Coronavirus (2019-nCov) IgM/IgG Antibody Combo Test Kit | see IFU | n/a | n/a | Hangzhou Laihe Biotech Co Ltd (China) distributed by Unisources Group LLC. | IgG/IgM | see IFU | see IFU | see IFU | IFU update | US FDA EUA |
| | 2019-nCoV/COVID-19 IgG/IgM Rapid Test Device | see IFU | n/a | n/a | Hangzhou Realy Tech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA removed from US FDA EUA |
| GCCOV-402a | COVID-19 IgG/IgM Rapid Test Cassette | 25 | n/a | n/a | Healgen Scientific Limited Liability Company (United States Of America) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA TGA |
| | InnoScreen™ COVID-19 IgG/IgM Rapid Test | see IFU | n/a | n/a | Innovation Scientific Pty Ltd (Australia) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|--|-------------------------|-------------------------------------|-----|--|----------|---------------------------------|---------------------------------|---------------|----------|--------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| YF319C | 2019-nCov Ab Test (Colloidal Gold) | see IFU | n/a | n/a | Innovita (Tangshan) Biological Technology Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA TGA |
| | Orawell IgM/IgG Rapid Test | see IFU | n/a | n/a | Jiangsu Well Biotech Co., Ltd. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | INDICAID COVID-19 IgM/IgG Rapid Test rebranded by Phase Scientific International Limited | see IFU | n/a | n/a | Jiangsu Well Biotech Co., Ltd. distributed by Phase Scientific International Limited | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | Rapid COVID-19 IgM/IgG Combo Test Kit | see IFU | n/a | n/a | Megna Health, Inc. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | COVID-19 IgG/IgM Rapid Test Kit | see IFU | n/a | n/a | Nantong Egens Biotechnology Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA removed from US FDA EUA |
| | COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Newscen Coast Bio-Pharmaceutical Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA removed from US FDA EUA |
| 8075 | ADEXUSDx COVID-19 Test | 50 | n/a | n/a | NOWDiagnostics, Inc | Total Ig | see IFU | see IFU | see IFU | | US FDA EUA |
| NBPC-0007 | MidaSpot™ COVID-19 Antibody Combo Detection Kit | 25 | n/a | n/a | Nirmidas Biotech, Inc | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|---|-------------------------|-------------------------------------|-----|--|---------|---------------------------------|---------------------------------|---------------|----------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| NBPC-0001-xx | Nirmidas COVID-19 (SARS-CoV-2) IgM/IgG Antibody Detection Kit | see IFU | n/a | n/a | Nirmidas Biotech, Inc | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | NTBIO One Step Rapid Test - COVID-19 IgG/IgM Antibody Test | see IFU | n/a | n/a | NTBIO Diagnostics Inc (Canada) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| | SARS-CoV-2 IgM/IgG Antibody Rapid Test | see IFU | n/a | n/a | Qingdao Hightop Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| CD-COV19CW/102223/102224 | Sienna-Clarity COVIBLOCK COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Salofa Oy | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| COGT025E, COGT005E | SGTi-flex COVID-19 IgG | 25T/kit 5T/kit | n/a | n/a | Sugentech, Inc | IgG | see IFU | see IFU | see IFU | | US FDA EUA |
| 20010 | TBG SARS-CoV-2 IgG / IgM Rapid Test Kit | see IFU | n/a | n/a | TBG Biotechnology Corp. | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| VID35-08-011 VID35-08-012 VID35-08-013 VID35-08-014 | VivaDiag™ COVID-19 IgM/IgG Rapid Test | 40T/kit | n/a | n/a | VivaCheck Biotech (Hangzhou) Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| | SARS-CoV-2 IgM/IgG Antibody Test Kit | see IFU | n/a | n/a | Wuhan EasyDiagnosis Biomedicine Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |

| SARS-CoV-2 Antibody Rapid Diagnostic Tests | | | | | | | | | | | |
|--|---|-------------------------|-------------------------------------|-----|---|---------|---------------------------------|---------------------------------|---------------|----------|-------------------------------|
| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
| | BIOTIME SARS-CoV-2 IgG/IgM Rapid Qualitative Test | 25T/kit | n/a | n/a | Xiamen Biotime Biotechnology Co., Ltd. (China) distributed by O’Neill Medical LLC, HORIBA instrument incorporated, Lifesaving Global LLC, THE RUHOF CORPORATION, Marlan’s Group Us Inc, and JEMF Pharma | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| | rebranded by Telepoint Medical Services, LLC | 25T/kit | n/a | n/a | Xiamen Biotime Biotechnology Co., Ltd. (China) distributed by Telepoint Medical Services, LLC | IgG/IgM | see IFU | see IFU | see IFU | | US FDA EUA |
| ☐ | BIOTIME SARS-CoV-2 IgG/IgM Rapid Qualitative Test | see IFU | n/a | n/a | Xiamen Biotime Biotechnology Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| ☐ GCCOV-402a | COVID-19 IgG/IgM Rapid Test Cassette | see IFU | n/a | n/a | Zhejiang Orient Gene Biotech Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| | COVID-19 IgG/IgM Rapid Test Kit | see IFU | n/a | n/a | Zhongshan Chuangyi Biochemical Engineering Co Ltd (China) | IgG/IgM | see IFU | see IFU | see IFU | | TGA |
| N/A- NOT APPLICABLE | | | | | | | | | | | |
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List of SARS-CoV-2 Diagnostic test kits and equipments eligible for procurement according to Board Decision on Additional Support for Country Responses to COVID-19 (GF/B42/EDP11)

A new technology for COVID-19 detection has become available that is much simpler and faster to perform than currently-recommended nucleic acid amplification tests (NAAT), like PCR. This method relies on direct detection of SARS-CoV-2 viral proteins in nasal swabs and other respiratory secretions using a lateral flow immunoassay (also called an RDT) that gives results in < 30 minutes. Though these antigen detection RDTs (Ag-RDTs) are substantially less sensitive than NAAT, they offer the possibility of rapid, inexpensive and early detection of the most infectious COVID-19 cases in appropriate settings. For more detailed technical advise, please, consult the WHO Interim guidance available at: <https://www.who.int/publications/i/item/antigen-detection-in-the-diagnosis-of-sars-cov-2infection-using-rapid-immunoassays>
https://www.who.int/publications/i/item/WHO-2019-nCoV-Ag-RDTs-Self_testing-2022.1

SARS-CoV-2 Nucleic Acid Amplification Tests for SELF TESTING
(Important note: Antigen tests do detect the omicron variant but may have reduced sensitivity and are therefore currently under risk assessment)

| Manufacturer Product Catalogue number | Product Name (IVD product) | Number of tests per kit | Platform (Calibrator and Equipment) | Manufacturer | Analyte | Anticipated Shelf life (months) | Recommended storage temperature | Specimen type | Comments | Eligibility WHO EUL or others |
|---------------------------------------|-------------------------------------|-------------------------|-------------------------------------|---------------------------------|---------|-----------------------------------|---------------------------------|---------------|--|---|
| 810-00012 Rev B | BinaxNOW COVID-19 Ag Card | 1Tests/kit | Metrix Reader | Aptitude Medical Systems | | see further product documentation | see IFU | see IFU | Reader required | US FDA EUA |
| 810055970056 | Lucira COVID-19 All-In-One Test Kit | 24T/kit | Disposable Lucira Device | Lucira Health, Inc. (by Pfizer) | N gene | see further product documentation | see IFU | see IFU | Under risk assessment for Omicron variant | US FDA EUA Health Canada/Interim Order |
| | Lucira CHECK-IT COVID-19 Test Kit | | Disposable Lucira Device | Lucira Health, Inc. | N gene | 18 months | see IFU | see IFU | For consumables and details of componants refer to IFU | US FDA EUA |

N/A- NOT APPLICABLE

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