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DIRECTORATE-GENERAL FOR HEALTH AND FOOD SAFETY

Public health
Health Security

EU HEALTH PREPAREDNESS

EU Common list of COVID-19 antigen tests

Agreed by the Health Security Committee

Last update: 22 July 2022

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1. Introduction

This document builds on the Council Recommendation of 21 January 2021¹, for which EU Member States unanimously agreed to set a common framework for the use and validation of antigen tests and the mutual recognition of COVID-19 test results across the EU. The Council Recommendation called for a common framework that consists of COVID-19 antigen tests that carry CE marking and that meet defined minimum performance requirements. Moreover, the devices should have been validated through an independent study carried out in at least one EU Member State, their use should be considered appropriate in the context of the COVID-19 pandemic, and their use should be in line with countries' national testing strategies.

COVID-19 rapid antigen tests

On 17 February 2021, and on the basis of the Council Recommendation of 21 January 2021, the Health Security Committee agreed on a **common list of COVID-19 rapid antigen tests**. Examples of rapid antigen tests are lateral flow immunoassays that give results in < 30 minutes.

The devices included in the EU common list of COVID-19 rapid antigen tests were meeting the criteria as defined by the Council Recommendation of 21 January 2021 as well as further criteria agreed by the Health Security Committee on 21 September 2021 (see 2.2). A dedicated Technical Working Group on COVID-19 diagnostic tests² was set up by Health Security Committee with the objective to assess proposals submitted by countries and manufacturers for new devices to be included in the EU common list against these agreed criteria.

Since its first publication in February 2021, the EU common list was regularly updated, taking into account the results of new validation studies and new rapid antigen test devices entering the market, as well as epidemiological developments and the emergence of SARS-CoV-2 variants.

EU common list of COVID-19 rapid antigen tests

- Agreed by the Health Security Committee on 17 February 2021
- First update: 10 May 2021; Second update: 16 June 2021; Third update: 7 July 2021; Fourth update: 14 July 2021; Fifth update: 23 July 2021; Sixth update: 20 October 2021; Seventh update: 10 November 2021; Eighth update: 8 December 2021; Ninth update: 21 December 2021; Tenth update: 21 January 2022; Eleventh update: 10 February 2022; Twelfth update: 4 March 2022; Thirteenth update: 8 April 2022; Fourteenth update: 6 May 2022; Fifteenth update: 10 June 2022.

As of 1 July 2021, and as determined by Regulation (EU) 2021/953 on the **EU Digital COVID Certificate**, all devices included in the EU common list of COVID-19 antigen tests and carried out by health professionals or by skilled testing personnel, could be used by EU Member States to issue EU Digital COVID test certificates. Moreover, as of 22 February 2022³, following a positive result of a COVID-19 rapid antigen test included in the EU common list and carried out

¹ Council Recommendation of 21 January 2021 on a common framework for the use and validation of rapid antigen tests and the mutual recognition of COVID-19 test results in the EU (OJ C 24, 22.1.2021, p.1).

² https://health.ec.europa.eu/health-security-and-infectious-diseases/crisis-management/covid-19-diagnostic-tests_en

³ Regulation (EU) 2021/953 of the European Parliament and of the Council of 14 June 2021 on a framework for the issuance, verification and acceptance of interoperable COVID-19 vaccination, test and recovery certificates (EU Digital COVID Certificate) to facilitate free movement during the COVID-19 pandemic (OJ L 211, 15.6.2021, p. 1–22).

by health professionals or by skilled testing personnel, it also became possible for EU Member States to issue EU Digital COVID recovery certificates. These certificates could be issued retroactively, based on rapid antigen tests carried out from 1 October 2021.

COVID-19 laboratory-based antigenic assays

In addition to the COVID-19 rapid antigen tests, since October 2021, the Health Security Committee has also agreed on a list of **mutually recognised COVID-19 laboratory-based antigenic assays**, such as enzyme-linked immunosorbent assays or automated immunoassays for the detection of antigens. These devices, meeting the same criteria as the COVID-19 rapid antigen tests, were until 30 June 2022 not eligible for the issuance of EU Digital COVID test and/or recovery certificates.

List of mutually recognised COVID-19 laboratory-based antigenic assays

- Agreed by the Health Security Committee on 20 October 2021
- First update: 10 February 2022; Second update: 8 April 2022; Third update: 10 June 2022.

The EU common list of COVID-19 antigen tests

As of 1 July 2022, as determined by Regulation (EU) 2022/1034⁴, both COVID-19 rapid antigen tests and COVID-19 laboratory-based antigenic assays are eligible for issuing EU Digital COVID test and recovery certificates. Therefore, on 22 July 2022, the Health Security Committee agreed that the EU common list of COVID-19 rapid antigen tests and the list of mutually recognised COVID-19 laboratory-based antigenic assays should be merged into one list: **the EU common list of COVID-19 antigen tests** (see Annex I).

All devices included in the EU common list of COVID-19 antigen tests are meeting the criteria as defined by the Council Recommendation of 21 January 2021 and further criteria agreed by the Health Security Committee on 21 September 2021 (see section 2.2.). All devices included in the EU common list of COVID-19 antigen tests and carried out by health professionals or by skilled testing personnel can be used by EU Member States to issue EU Digital COVID test and recovery certificates.

EU common list of COVID-19 antigen tests

- Agreed by the Health Security Committee on 22 July 2022.

As stipulated in point 15 of the Council Recommendation of 21 January 2021, Member States will agree on a selection of antigen tests of which they will mutually recognise the test results for public health measures. The Health Security Committee agrees that, considering that *all* of the antigen tests, both rapid antigen tests and laboratory-based antigenic assays, included in the EU common list are eligible for issuing EU Digital COVID test and recovery certificates, the entire list is considered to consist of antigen tests of which Member States mutually recognise the test results for public health measures.

⁴ Regulation (EU) 2022/1034 of the European Parliament and of the Council of 29 June 2022 amending Regulation (EU) 2021/953 of the European Parliament and of the Council of 14 June 2021 on a framework for the issuance, verification and acceptance of interoperable COVID-19 vaccination, test and recovery certificates (EU Digital COVID Certificate) to facilitate free movement during the COVID-19 pandemic (OJ L 173, 30.6.2022, p. 37–45).

2. The EU common list of COVID-19 antigen tests

2.1 Category A and Category B devices

The EU common list of COVID-19 antigen tests has been split up in two categories:

- **Category A:** Antigen tests for which their performance has been evaluated through prospective clinical field studies and that meet the criteria agreed on 21 September 2021 (see section 2.2) have been placed under the “A-category” of the EU common list. **Category A.1** sets out the eligible COVID-19 rapid antigen tests and **Category A.2** sets out the eligible COVID-19 laboratory-based antigenic assays.
- **Category B:** Antigen tests for which their performance has been evaluated through retrospective in vitro studies and that meet the criteria agreed on 21 September 2021 (see section 2.2) have been placed under the “B-category” of the EU common list. **Category B.1** sets out the eligible COVID-19 rapid antigen tests and **Category B.2** sets out the eligible COVID-19 laboratory-based antigenic assays.

EU Member States are strongly encouraged to use, in particular, antigen tests included under Category A of the EU common list for the issuance of EU Digital COVID certificates.

Secondly, EU Member States should pay particular attention to the issuance of EU Digital COVID recovery certificates based on the result of devices listed under Category B and that have solely been evaluated by the Paul-Ehrlich-Institut (PEI) in Germany, as only the sensitivity of these antigen tests has been evaluated.

Thirdly, EU Member States are strongly encouraged to ensure that only test results from the evaluated specimen type(s) as indicated for Category A devices are used for the issuance of EU Digital COVID test and recovery certificates. As regards the Category B devices, in general, retrospective in vitro studies do not aim to evaluate the clinical performance of an antigen test based on a specific specimen type. Therefore, the clinical performance of devices listed under Category B cannot be linked to a specific specimen type, which should be taken into consideration by EU Member States when using these antigen tests for the issuance of EU Digital COVID certificates.

2.2 Criteria to be met

Based on a proposal by their Technical Working Group and taking into account the criteria presented by the Council Recommendation of 21 January 2021, the following section sets out the scope, definitions and criteria that were agreed by the Health Security Committee agreed on 21 September 2021 and that should be met by devices in order to be included in the EU common list of COVID-19 antigen tests.

The Technical Working Group of the Health Security Committee monitors technical and epidemiological developments in the field of antigen testing on a continuous basis and will, if deemed necessary, reconsider the scope, definitions and criteria to be met by devices included in the EU common list. Particular attention will be paid to breakthrough infections among vaccinated individuals and the possible impact of such cases on the clinical performance of

antigen tests, as well as the performance of antigen tests in the context of emerging SARS-CoV-2 variants. Moreover, the ongoing work by the In Vitro Diagnostics Working Group of the Medical Device Coordination Group regarding guidance on the performance of COVID-19 tests in the context of CE-marking and common specifications under Article 9 of Regulation (EU) 2017/746⁵ will be taken into account. If considered relevant, a proposal for an update of the below agreements will be put forward by the Technical Working Group to the Health Security Committee.

Agreed scope of the EU common list of COVID-19 antigen tests:

- Only antigen tests that carry CE marking are included in the EU common list of COVID-19 antigen tests.
- The EU common list includes antigen tests that are used in practice in and that have been validated by at least one of the 27 EU Member States⁶.
- The EU common list includes antigen tests for which their clinical performance was measured based on samples collected from nasal, oropharyngeal or nasopharyngeal specimens and that meet the criteria as further specified below.
- Antigen tests that are using a mix of different sampling materials (i.e. nasal, oropharyngeal and/or nasopharyngeal swabs as well as other specimen types such as saliva) can be included in the EU common list.
- In case antigen tests are based on using multiple sampling materials, each specimen type should be evaluated separately and the results and data of validation studies should thus be presented per specimen type. The EU common list indicates for devices evaluated by prospective field studies, which of the specimen types have been evaluated and which of the specimen types meet the agreed criteria. Note that only the results of validation studies based on nasal, oropharyngeal and/or nasopharyngeal swabs of such devices will be reviewed by the Technical Working Group and assessed against the specified criteria.
- Only test results based on nasal, oropharyngeal and/or nasopharyngeal specimens should be valid for the issuance of test certificates for the EU Digital COVID Certificate.
- Multiplex antigen assays that simultaneously detect multiple pathogens – e.g. SARS-CoV-2 and Influenza A/B - can be included in the EU common list, but only their performance regarding their detection of SARS-CoV-2 will be assessed.

The EU common list of COVID-19 antigen tests does **NOT** include, and therefore EU Digital COVID certificates cannot be issued based on a test result from:

- Antigen tests that are solely based on sampling materials other than nasal, oropharyngeal or nasopharyngeal specimens, such as saliva, sputum, blood and/or faeces. This is in line with current evidence and the technical recommendations provided by the European Centre for Disease Prevention and Control (ECDC)⁷.

⁵ The Medical Device Coordination Group is set up according to Art. 103 of Regulation (EU) 2017/745 and Art. 98 of Regulation (EU) 2017/746. This group is also responsible for overseeing the implementation of Directive 98/79/EC. See also Register of Commission Expert Groups and Other Similar Entities, code number X03565, and its subgroups.

⁶ https://european-union.europa.eu/principles-countries-history/country-profiles_en.

⁷ <https://www.ecdc.europa.eu/sites/default/files/documents/covid-19-use-saliva-sample-material-testing.pdf>.

- Antigen self-tests, including antigen self-testing monitored by health professionals or by skilled testing personnel (either on site or remotely). The EU common list only includes antigen tests that are conducted by trained healthcare personnel or trained operators where appropriate (in line with Commission Recommendation (EU) 2020/1743 of 18 November 2020).
- Pooled antigen tests, which involve mixing of multiple samples together in a batch or pooled sample for testing.
- COVID-19 antibody tests, nor does the Technical Working Group assess the performance of these tests against the agreed criteria.

Agreed criteria and definitions of an independent validation study:

The clinical performance of antigen tests included in the EU common list should have been evaluated by an independent validation study meeting the following criteria and definitions:

- The validation study should be performed by an independent laboratory, which is a laboratory not owned nor operated by the manufacturer or sponsor of the test, and which is not related to the manufacturer/sponsor of the test by ownership, familial relationships, nor contractual or other relationships that result in the laboratory being controlled by or being under the common control of the manufacturer/sponsor of the test.
- The independent validation study should have been carried out in at least one of the 27 EU Member States⁸, and be performed objectively and in the public interest.
- The independent validation study may involve collaborations with or may involve funding by private entities, however, there is always a public body involved from an EU Member State.
- The independent validation study should preferably be based on a **prospective clinical field study** design, testing *unselected* symptomatic and asymptomatic participants for SARS-CoV-2 infection.
- “Unselected” means no prior knowledge of SARS-CoV-2 diagnosis (e.g. determined by PCR); inclusion is allowed based on general possible COVID-like symptoms (or close contact with COVID-19 cases); and exclusion is allowed of children (e.g. <16 years) or for medical ethical permission reasons.
- The performance of antigen tests can also be evaluated based on a **retrospective in vitro study design**, testing the clinical performance by using SARS-CoV-2 reference panels.

Agreed clinical performance criteria for independent validation studies:

Prospective clinical field studies:

- A sensitivity over 80% when testing unselected symptomatic participants within the first seven days after symptom onset or asymptomatic participants, where the diagnosis is confirmed by RT-PCR in independent field studies, will be accepted.

⁸ https://european-union.europa.eu/principles-countries-history/country-profiles_en.

OR

In independent evaluations of unselected participants, assays should have a sensitivity of 90% or greater for subjects with a $Ct \leq 25$.

- The study population shall be clearly defined stating the inclusion criteria of participants (symptomatic individuals, close contacts or asymptomatic individuals without known exposure). Ideally, the sensitivity for each group should be discernible from the report. The RT-PCR protocol and the distribution of Ct values should be described. Samples should represent naturally occurring viral loads.
- Target population should be based on at least 100 fresh RT-PCR positive samples and at least 300 fresh RT-PCR negative samples. Each specimen type should be evaluated separately.
- In case of multiple smaller prospective clinical field studies that do not meet the minimum number of positive and/or negative samples separately but that do meet all the other criteria as agreed by the Technical Working Group, the number of samples may be combined, provided that the different studies applied the same or similar methodologies, are all carried out in EU Member States, and that sufficient details are provided on their study design.
- Assays should have a specificity over 98%.
- In line with the *MDCG Guidance on performance evaluation of SARS-CoV-2 in vitro diagnostic medical devices*⁹, preference is given to samples being compared against RT-PCR results on nasopharyngeal swabs. However, in independent validation studies, samples can also be compared against RT-PCR results on oropharyngeal or nasal swabs if reasoning is provided (e.g. when assessing the clinical performance of antigen tests among children).

Retrospective in vitro studies:

- A sensitivity over 80% when testing all specimen in the reference panel will be accepted;

OR

Assays should have a sensitivity of 90% or greater for subjects with a $Ct < 25$.

- The composition of the reference panel should be as follows:
 - A panel of at least 50 (pooled) clinical specimens that cover naturally occurring viral loads with SARS-CoV-2 concentration ranging from approximately 1.1×10^9 to 4.2×10^2 genome copies per mL of specimen and Ct values between 17 and 36.
 - The whole evaluation panel should be subdivided into three subgroups: panel members, which are characterized by:
 - **Very high viral load:** Ct value 17-25; 35% (+/- 2%) of the total number of (pooled) clinical specimens);
 - **High viral load** (Ct value 25-30; 45% (+/- 2%) of the total number of (pooled) clinical specimens); and
 - **Moderate viral load** (Ct value 30-36; 20% (+/- 2%) of the total number of (pooled) clinical specimens).

⁹ https://ec.europa.eu/health/sites/default/files/md_sector/docs/mdcg_2021-21_en.pdf

- For each pool up to ten clinical respiratory specimens (nasopharyngeal/oropharyngeal) obtained for routine diagnostics with different virus loads may be used. The sample volume per panel member should be sufficient to allow comparative evaluation with different tests included in the evaluation.
 - RT-PCR needs to be applied to determine the RNA load per pool.
 - Ethical approval by an institutional review board is mandatory.
 - For each antigen test and panel member, a pre-defined aliquot needs to be completely absorbed using the specimen collection device, e.g. swab, provided with the respective test.
 - Further steps needs to be strictly performed following the respective instructions for use (IFU).
 - The stability of the panel (antigen) must be considered throughout the preparation of the panel and the workflow up to the test.
- Assays should have a specificity over 98%, as measured through the retrospective in vitro evaluation study or as specified by the manufacturer in the IFU.
 - In line with the *MDCG Guidance on performance evaluation of SARS-CoV-2 in vitro diagnostic medical devices*¹¹, preference is given to samples being compared against RT-PCR results on nasopharyngeal swabs. However, in independent validation studies, samples can also be compared against RT-PCR results on oropharyngeal or nasal swabs if reasoning is provided (e.g. when assessing the clinical performance of antigen tests among children).

2.3 Process to include devices in the EU common list

As called for by the Council Recommendation of 21 January 2021, the EU common list of COVID-19 antigen tests should feed into the “COVID-19 In Vitro Diagnostic Devices and Test Methods Database”¹⁰, hosted by the Joint Research Centre (JRC). Therefore, as of May 2021, it is possible for EU Member States as well as manufacturers of COVID-19 antigen tests to put forward proposals of antigen tests to be included in the EU common list by submitting the relevant information to the COVID-19 In Vitro Diagnostic Devices and Test Methods Database.

After following the correct procedures and once verified against the source provider, the proposals will be forwarded to the Technical Working Group of the Health Security Committee for evaluation and review. Note that the submission of an application does not result in an immediate review, as the Technical Working Group assesses batches of proposals that have been submitted during a pre-defined period.

2.4 Updating of the EU common list

The Technical Working Group of the Health Security Committee meets, on average, once a month, during which the next batch of proposals by manufacturers and countries is reviewed and discussed. Based on these discussions, a proposal for a possible next update of the EU common list is forwarded to the Health Security Committee for review and formal agreement.

¹⁰ <https://covid-19-diagnostics.jrc.ec.europa.eu/devices>.

Once the Health Security Committee has agreed with the update, it will be published on the Commission's website and the COVID-19 In Vitro Diagnostic Devices and Test Methods Database will be updated accordingly.

In parallel to the latest update of the EU common list, the Health Security Committee also agrees with the publication of an updated **addendum**. This document provides further background information to the decisions taken by the Technical Working Group in the context of the latest update to the common list of COVID-19 antigen tests. It provides manufacturers and EU Member States with an overview of the devices of which their inclusion in the EU common list was rejected as well as the proposals that are still under review.

Apart from the inclusion of new devices, the Technical Working Group may also decide that certain antigen tests should be removed from the EU common list. For example, this may happen in case new validation results are published, showing that the device no longer meets the agreed criteria or, in case a new variant emerges that affects the clinical performance of certain antigen tests.

Grace period

As of 1 January 2022, a grace period of 4 weeks applies whenever updates are made to Annex I. The grace period applies to both the inclusion of new devices as well as the removal of antigen tests that are included in the EU common list or COVID-19 antigen tests.

3. Background information

COVID-19 antigen tests

Robust and targeted testing strategies are an essential aspect of preparedness and response to the COVID-19 pandemic. Representative testing strategies provide useful indications on the epidemiological trends and of the intensity of community transmission, the impact of severe disease and on vaccine effectiveness. Moreover, they are a prerequisite to adequate contact tracing to limit the spread through prompt isolation. Also in the context of the circulation of SARS-CoV-2 variants of concern, testing is pivotal to ensuring representative and targeted genomic sequencing efforts.

While the reverse transcription real-time polymerase chain reaction (RT-PCR) assay, which is a nucleic acid amplification test (NAAT), remains the 'gold standard' for COVID-19 diagnosis, antigen tests, which detect the presence of viral proteins (antigens), are being used by Member States as a way of further strengthening overall testing capacity, particularly in case of limited NAAT capacities or where clinical needs require faster testing turnaround times.

The Health Security Committee

The Health Security Committee, set up in 2001, is mandated to reinforce the coordination and sharing of best practice and information on national health security activities as well as the coordination of national responses to serious cross border threats to health, including events

declared a public health emergency of international concern by World Health Organization in accordance with the International Health Regulations¹¹.

On 17 September 2020, the Health Security Committee agreed on Recommendations for a common EU testing approach for COVID-19¹², which included Member States' first experiences with antigen tests and their deliberations concerning the settings and situations in which these tests should be used. Since then, the HSC has been discussing the use of antigen tests in great depth, and has brought together a wealth of (technical) information on the types of tests used in European countries and the conditions applied.

In May 2021, the Health Security Committee set up a **Technical Working Group on COVID-19 Diagnostic Tests**, consisting of technical experts from EU and EEA Member States, as well as representatives from the Directorate-General for Health and Food Safety (DG SANTE), the Commission's Joint Research Centre (JRC) and the European Centre for Disease Prevention and Control (ECDC). The Technical Working Group is responsible for reviewing the information submitted by countries and manufacturers, taking into account the latest result of independent validation studies and country practices and experiences. Based on this, the Technical Working Group presents proposals for further updates to the common list of COVID-19 antigen tests to the Health Security Committee for agreement.

¹¹ https://ec.europa.eu/health/health-security-and-infectious-diseases/preparedness-and-response/health-security-committee-hsc_en.

¹² https://ec.europa.eu/health/sites/health/files/preparedness_response/docs/common_testingapproach_covid-19_en.pdf

ANNEX I: EU common list of COVID-19 antigen tests ^{13, 14}

Disclaimer: The Technical Working Group strongly recommends that antigen tests are primarily used for preliminary testing for SARS-CoV-2 infection in symptomatic patients, and notes that antigen tests should in particular be used in the specific contexts and circumstances referred to by the Commission Recommendation (EU) 2020/1743 and the updated technical report by ECDC on 26 October 2021. The content of the EU common list is based on the clinical performance data and information that is available at this moment in time. The Technical Working Group stresses that the clinical performance data of devices included in the EU common list, resulting from independent validation studies meeting the agreed criteria, cannot be directly compared as absolute numbers.

Category A: COVID-19 antigen tests evaluated by prospective clinical field studies

EU Member States are strongly encouraged to use, in particular, the antigen tests included under “Category A” of the EU common list for the issuance of EU Digital COVID certificates. The clinical performance of these devices – for the specimen type as indicated in the corresponding column - has been evaluated by (at least) one prospective clinical field study meeting the criteria and definitions as agreed by the Health Security Committee on 21 September 2021.

EU Member States are strongly encouraged to ensure that only test results from the evaluated specimen type(s) are used to issue EU Digital COVID certificates.

Devices highlighted in blue are identical in design and construction but are, for example, branded or distributed under a different name. The results of validation studies may be transferred between devices that are identical in design and construction.

CATEGORY A.1: COVID-19 RAPID ANTIGEN TESTS

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since <i>D/M/Y</i>
1833	AS-COV-008, AS-COV-009	AAZ-LMB	COVID-VIRO®	Prospective clinical field study Prospective study carried out in the “Centre Hospitalier d’Orléans” on nasopharyngeal swabs, simultaneously tested by RT PCR:	Nasopharyngeal	Nasal	Nucleocapsid protein	10/05/2021

¹³ This is the list of COVID-19 antigen tests as referred to in Article 3 of the Regulation (EU) 2022/1034 of the European Parliament and of the Council of 29 June 2022, amending Regulation (EU) 2021/953 of the European Parliament and of the Council of 14 June 2021 on a framework for the issuance, verification and acceptance of interoperable COVID-19 vaccination, test and recovery certificates (EU Digital COVID Certificate) to facilitate free movement during the COVID-19 pandemic, OJ L 173, 30.6.2022, p. 37–45.

¹⁴ The Medical Device Coordination Group [Guidance on performance evaluation of SARS-CoV-2 in vitro diagnostic medical devices](#), which will form the basis for common specifications to be adopted according to Article 9 of Regulation (EU) 2017/746, has been considered by the Technical Working Group for the development of the EU common list of COVID-19 antigen tests.

¹⁵ As registered in and used by the JRC database; see: <https://covid-19-diagnostics.jrc.ec.europa.eu/>.

¹⁶ The reference number is the identification number issued by the manufacturer to identify the device. It is usually included in the device’s labelling, instructions for use and/or declaration of conformity, and often preceded by the symbol ‘REF’. Synonyms for reference numbers are: catalogue numbers, commercial product codes or reorder numbers. The reference number may vary in different markets. The REF number included in the EU common list may be followed by “{...}”, which means this is generic number that will, in practice, be followed by further references, depending on the packaging, box, national market, etc. When a reference number is not issued, the device is typically identified by its commercial product name.

¹⁷ Identical to what is included in the Instructions For Use (IFU) and/or labelling of the antigen test.

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				sensitivity <7 days after onset of symptoms: 94.7%, specificity: 100%.				
1232	41FK10	Abbott Rapid Diagnostics (manufacturer)	Panbio™ COVID-19 Ag Rapid Test	<p>Prospective clinical field study Study enrolling 1367 and 208 subjects in Utrecht (NL) and Aruba, respectively. NP swabs. Specificity was 100% (95%CI: 99.7–100%) in both settings. Test sensitivity was 72.6% (95%CI: 64.5–79.9%) in the Netherlands and 81.0% (95% CI: 69.0–89.8%) in Aruba. Restricting RT-qPCR test positivity to Ct-values <32 yielded test sensitivities of 95.2% (95%CI: 89.3–98.5%) in Utrecht and 98.0% (95%CI: 89.2–99.95%) in Aruba. Source.</p> <p>FIND prospective evaluation study Germany (10 Dec 2020): 1108 samples, NP swab. Clinical sensitivities: Days ≤ 7: 90.8%; Ct ≤ 33: 88.3%; Ct ≤ 25: 95.8%. Clinical specificity: 99.9%. Source.</p> <p>Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.8%</i>.</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	17/02/2021
1457	L031-11815, L031-125A5	Acon Biotech (Hangzhou) Co., Ltd (manufacturer)	Flowflex SARS-CoV-2 Antigen Rapid Test	<p>Prospective clinical field study Independent prospective study carried out by University Magna Graecia of Catanzaro, Italy. Unselected nasal samples: 144 positive and 342 negative samples. Sensitivity: 97.22%, specificity: 99.71%. Unselected NP samples: 145 positive and 332 negative samples. Sensitivity: 96.55%, specificity: 100%.</p> <p>Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.54%</i>.</p>	Nasal, Nasopharyngeal	-	Nucleocapsid protein	14/07/2021
2079		ArcDia International Ltd	mariPOC Quick Flu+	<p>Prospective clinical field study Clinical performance of the test was evaluated in Finland against qRT-PCR with NP swab specimens collected from patients suspected of acute SARS-CoV-2 infection. Sensitivity of the</p>	Nasopharyngeal	-	Nucleocapsid protein	14/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				<p>mariPOC test was 100.0% (13/13) directly from swab specimens and 84.4% (38/45) from swab specimens in undefined transport mediums. Specificity was 100.0% (201/201). Source.</p> <p>Prospective clinical field study Clinical performance of the test was evaluated in Finland against RT-PCR with specimens from 962 symptomatic and asymptomatic individuals. Among the symptomatic subjects, overall sensitivity was 82.5% (33/40), which increased to 97.1% (33/34) in samples with a Ct value <30. The specificity was 100% (916/916). Source.</p>				
2078	1184M	ArcDia International Ltd	mariPOC Respi+	<p>Prospective clinical field study Clinical performance of the test was evaluated in Finland against qRT-PCR with NP swab specimens collected from patients suspected of acute SARS-CoV-2 infection. Sensitivity of the mariPOC test was 100.0% (13/13) directly from swab specimens and 84.4% (38/45) from swab specimens in undefined transport mediums. Specificity was 100.0% (201/201). Source.</p> <p>Prospective clinical field study Clinical performance of the test was evaluated in Finland against RT-PCR with specimens from 962 symptomatic and asymptomatic individuals. Among the symptomatic subjects, overall sensitivity was 82.5% (33/40), which increased to 97.1% (33/34) in samples with a Ct value <30. The specificity was 100% (916/916). Source.</p>	Nasopharyngeal	-	Nucleocapsid protein	14/07/2021
768	1204S	ArcDia International Ltd	mariPOC SARS-CoV-2	<p>Prospective clinical field study Clinical performance of the test was evaluated in Finland against qRT-PCR with NP swab specimens collected from patients suspected of acute SARS-CoV-2 infection. Sensitivity of the mariPOC test was 100.0% (13/13) directly from swab specimens and 84.4% (38/45) from swab specimens in undefined transport mediums. Specificity was 100.0% (201/201). Source.</p> <p>Prospective clinical field study Clinical performance of the test was evaluated</p>	Nasopharyngeal	-	Nucleocapsid protein	10/05/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				in Finland against RT-PCR with specimens from 962 symptomatic and asymptomatic individuals. Among the symptomatic subjects, overall sensitivity was 82.5% (33/40), which increased to 97.1% (33/34) in samples with a Ct value <30. The specificity was 100% (916/916). Source .				
2282	256091, 256113, 256114	Becton Dickinson (manufacturer)	BD Kit for Rapid Detection of SARS-CoV-2	Prospective clinical field study Study in four Spanish hospitals (n = 476); 108 positive samples, 368 negative samples. Sensitivity: 92%, specificity: 98.6%.	Nasal	-	Nucleocapsid protein	10/11/2021
1065	256089		BD Veritor™ System for Rapid Detection of SARS CoV 2	Prospective clinical field study Independent study in the Netherlands including symptomatic individuals (n=979, PCR positive n=161). Sampling: Nasal mid-turbinate and OP swabs. Sensitivity overall: 79.5%; Sensitivity Ct<30: 93.2%; Specificity overall: 99.8%.				07/07/2021
1778	601460, 601540	Beijing Kewei Clinical Diagnostic Reagent Inc	COVID19 Antigen Rapid Test Kit	Prospective clinical field study Study at the General Hospital Jesenice in Slovenia: 103 RT-PCR positives and 450 RT-PCR negative subjects; symptomatic patients only. Overall sensitivity: 91.26%; specificity: 99.33%. Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%</i> .	Nasal	-	Nucleocapsid protein	21/12/2021
1485	WJ-2950	Beijing Wantai Biological Pharmacy Enterprise Co., Ltd	Wantai SARS-CoV-2 Ag Rapid Test (colloidal gold)	Prospective clinical field study Independent prospective study by Public Health Institute Ostrava (Czechia), including nasopharyngeal swabs from unselected symptomatic and asymptomatic participants. Sensitivity 80.6%, specificity 98.5% on 155 positive and 325 negative samples against RT-PCR (N total = 480). Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.2%</i> .	Nasopharyngeal	Nasal	Nucleocapsid protein	14/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
3099		BioDetect (Xiamen) Biotechnology Co., Ltd.	RAPID SARS-COV-2 ANTIGEN TEST CARD	Prospective clinical field study Independent prospective study in Slovenia, General Hospital Jesenice. Nasal swab, symptomatic and asymptomatic patients. Positive samples: 200, negative samples: 404. Sensitivity: 90% (180/200), specificity: 99.75% (403/404).	Nasal	-	Nucleocapsid protein	22/07/2022
2519	KACOV19-5, KACOV19-25, KACOV19-25C	BIOLAN HEALTH, S.L.	COVID-19 Antigen Rapid Test (Colloidal Gold Method)	Prospective clinical field study Prospective study performed in Hospital Universitario de Cruces (Spain). Nasal specimen, 314 negative samples and 116 positive samples. Sensitivity 98.1% at Ct<25; overall sensitivity 81%; specificity 98.1%.	Nasal	-	Nucleocapsid protein	04/03/2022
2035	1-367-K020	BioMaxima SA	SARS-CoV-2 Ag Rapid Test	Prospective clinical field study Study in Poland performed on 480 samples of NP swabs taken from symptomatic patients and from asymptomatic people in contact with an infected person. Positive results were obtained in 205 patients and in the molecular test 213 people. Negative results were obtained in 275 people and in the molecular test 267 people. Diagnostic sensitivity: 93.43% (95% CI: 91.61%~97.19%) and diagnostic specificity: 97.75% (95% CI: 93.74%~98.92%). Source . Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%</i> .	Nasopharyngeal	-	Nucleocapsid protein	23/07/2021
2031	12015553	BIO-RAD	CORONAVIRUS AG RAPID TEST CASSETTE	Prospective clinical field study Study carried out in Spain; 96 positive samples and 269 negative samples. Sensitivity 94%. Specificity 99.2%. Prospective clinical field study Study carried out in Spain; nasopharyngeal swabs, sensitivity 98.3%; specificity 99.6% (119 positive samples, 746 negative samples). Prospective clinical field study Study carried out in Spain; nasal swabs, sensitivity 97.2%; specificity 100% (109 positive samples, 128 negative samples).	Nasopharyngeal, Nasal	-	Nucleocapsid protein	07/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2380	BSD_0503-10, BSD_0503-25	BioSpeedia International	COVID19Speed-Antigen Test BSD_0503	<p>Prospective clinical field study</p> <p>Independent prospective study by the University Hospital of Saint-Etienne (France): samples from unselected symptomatic and asymptomatic individuals (255 pos., 365 neg.), overall sensitivity: 95.29% (sensitivity Ct<25: 97.72%), specificity: 99.73%.</p>	Nasopharyngeal	-	Nucleocapsid protein	21/01/2022
1494	SW40010	BIOSYNEX SA	BIOSYNEX COVID-19 Ag+ BSS	<p>Prospective clinical field study</p> <p>Validation study carried out in France: 125 positive and 118 negative samples; sensitivity 96%, specificity: 99%.</p> <p>Prospective clinical field study</p> <p>Clinical study carried out in a public health hospital in France (centre cardiologique du Nord): sensitivity 100% (188/188), specificity 100% (313/313).</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	07/07/2021
1223	SW40006	BIOSYNEX SWISS S.A. (manufacturer)	BIOSYNEX COVID-19 Ag BSS	<p>Prospective clinical field study</p> <p>Independent field study in the Netherlands, involving mainly symptomatic individuals (n=568, PCR positive n=39), NP swab; sensitivity Ct ≤ 30: 96.0%, sensitivity Ct ≤ 25: 100%; specificity overall: 100%.</p> <p>Prospective clinical field study</p> <p>Independent field study in the Netherlands, symptomatic individuals (n=270, PCR positive n=17), NP+OP swab; sensitivity Ct ≤ 30: 94.1%, sensitivity Ct ≤ 25: 100%; specificity: 100%.</p> <p>Prospective clinical field study</p> <p>Prospective study in France, nasopharyngeal swabs (n=71/71): sensitivity 100% (45/45, specificity 100%.</p> <p>Prospective clinical field study</p> <p>Evaluation in Karolinska hospital (Sweden) of Lot 20100103. Patient samples; 95 PCR positive, 150 negative. Sensitivity 76%, specificity 96%. Sensitivity Ct<25 = 100%.</p> <p>Retrospective in vitro study</p> <p>Positive evaluation by Paul-Ehrlich-Institut (PEI)</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	17/02/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>				
1989	SMFP-71	Boditech Med Inc	AFIAS COVID-19 Ag	Prospective clinical field study Independent field study in the Netherlands in mild symptomatic (n= 427, PCR positive: 106); overall sensitivity: 81.1%, sensitivity Ct <30: 96.4%; specificity: 100%.	Nasopharyngeal	-	Nucleocapsid protein	23/07/2021
1647		CALTH Inc.	AllCheck COVID19 Ag	Prospective clinical field study Independent prospective study carried out in the laboratory of a public hospital in Italy. Sample size: 136 positive and 520 negative samples. Overall sensitivity: 95.6%; Sensitivity 98.9% for Ct ≤ 25; specificity: 100%.	Nasopharyngeal	-	Nucleocapsid protein	22/07/2022
2691		CALTH Inc.	AllCheck COVID19 Ag Nasal	Prospective clinical field study Independent prospective study carried out in the laboratory of a public hospital in Italy. Sample size: 130 positive and 530 negative samples. Overall sensitivity: 95.4% (124/130), specificity: 99.8% (519/530).	Nasal	-	Nucleocapsid protein	22/07/2022
1173	SC820001PC	CerTest Biotec	CerTest SARS-CoV-2 Card test	Prospective clinical field study Clinical study in Spain by Hospital Universitario Príncipe de Asturias (Madrid) during January/February 2021 and May 2022. Sample size: 98 positive and 242 negative samples; NP swabs. Overall sensitivity: 65% (92% for Ct ≤ 25), specificity: 98%. Prospective clinical field study Prospective study on NP samples by Laboratoire de Virologie CHU (Amiens, France) during October/November 2020. Sample size: 96 positive and 62 negative samples. Sensitivity 92% for Ct ≤ 25, specificity: 98%.	Nasopharyngeal	Nasal	Nucleocapsid protein	17/02/2021
1225	04 nCov	DDS DIAGNOSTIC	Test Rapid Covid-19 Antigen (tampon nazofaringian)	Prospective clinical field study Clinical study in Romania based on 228 positive samples and 597 negative samples. All the samples were confirmed using PCR (Applied Biosystems™ 7500 and SLAN®- 96P) and clinical symptoms. The relative sensitivity (nasopharyngeal Swab) was 99.56%, the relative specificity was 99.66%.	Nasopharyngeal	-	Nucleocapsid protein	10/05/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2147	231906	Fujirebio	ESPLINE SARS-CoV-2	FIND prospective evaluation study Germany (29 March 2021): 723 samples, NP swab. Sensitivities: Days < 7: 88.5%; Ct < 33: 87.8%; Ct < 25: 92.4%. Clinical specificity: 100%. Source . Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.13%</i> .	Nasopharyngeal	-	Nucleocapsid protein	07/07/2021
1144	643G	Green Cross Medical Science Corp.	GENEDIA W COVID-19 Ag	Prospective clinical field study Independent prospective study by University Hospital Sant'Andrea in Rome, Italy. Sample size: 117 positive and 400 negative samples, unselected population. Overall sensitivity: 82.05%, specificity: 100%	Nasopharyngeal	Anterior nasal	Nucleocapsid protein	10/05/2021
1747	JT04-20	Guangdong Hecin Scientific, Inc.	2019-nCoV Antigen Test Kit (colloidal gold method)	Prospective clinical field study Study in a public hospital in Slovenia with acute symptomatic patients, Nasal samples. Sample size: 103 positive, 450 negative samples. Sensitivity: 97.09% (100/103) and specificity: 99.78% (449/450).	Nasal	Nasopharyngeal	Nucleocapsid protein	10/05/2021
2302	FI-NCP-502	Hangzhou AllTest Biotech Co., Ltd	COVID-19 Antigen Test Cassette (Nasopharyngeal Swab) (FIA)	Prospective clinical field study Study in a public hospital in Slovenia, unselected patients, NP samples, 102 positive samples and 312 negative samples. Sensitivity: 95.1% and specificity: 100%.	Nasopharyngeal	-	Nucleocapsid protein	08/04/2022
1257	INCP-502	Hangzhou AllTest Biotech Co., Ltd	SARS-CoV-2 Antigen Rapid Test (COVID-19 Antigen Rapid Test) (Swab)	Prospective clinical field study Study at General Hospital Jesenice in Slovenia, unselected asymptomatic and symptomatic participants. Sample size: 127 positive, 316 negative. Sensitivity: 97.6% (124/127); specificity: 99.7% (315/316).	Nasopharyngeal	-	Nucleocapsid protein	10/05/2021
2319	CVAG4080A		CVAG4080A – GSD NovaGen SARS-CoV-2 Ag Rapid Test (NP Swab)					10/06/2022
1375	Z20401CE (rev17), Z20999CE (rev 16), Z20601CE (rev13)		DIALAB GmbH					DIAQUICK COVID -19 Ag Cassette

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1767	GCCOV-502a	Healgen Scientific (manufacturer)	Coronavirus Ag Rapid Test Cassette	<p>Prospective clinical field study</p> <p>Clinical field study in the Netherlands including symptomatic individuals (n=417, PCR positive n=70), NP swab; sensitivity overall: 75.7%, sensitivity Ct ≤ 30: 85.2%, sensitivity Ct ≤ 25: 90.7%; specificity: 100%.</p> <p>Prospective clinical field study</p> <p>Clinical field study in the Netherlands including symptomatic individuals (n=240, PCR positive n=21), NP+OP swab; sensitivity overall: 85.7%, sensitivity Ct ≤ 30: 89.5%, sensitivity Ct ≤ 25: 100%; specificity: 100%.</p>				17/02/2021
1218	GCCOV-502a	Siemens Healthineers (manufacturer)	CLINITEST Rapid COVID-19 Antigen Test	<p>Prospective clinical field study</p> <p>Clinical field study in the Netherlands including symptomatic individuals (n=94, PCR positive n=18), NP+OP swab in VTM; sensitivity overall: 90.0%, sensitivity Ct ≤ 30: 100%, sensitivity Ct ≤ 25: 100%; specificity: 97.3%.</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	17/02/2021
1343	GCCOV-502a	Zhejiang Orient Gene Biotech Co., Ltd (manufacturer)	Coronavirus Ag Rapid Test Cassette (Swab)	<p>Prospective clinical field study</p> <p>Independent prospective study in Spain: 192 positive and 258 negative samples (NP swab). Sensitivity: 93.3%, specificity: 99.2%, compared against NP PCR.</p> <p>Retrospective in vitro study</p> <p>Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%</i>.</p>				17/02/2021
1791	RTVI039AG	Immunospark s.r.l.	Rapid SARS-Cov2 Antigen Test	<p>Prospective clinical field study</p> <p>Study with unselected individuals (with delayed antigen testing), supervised by a public university in Italy. Sample size (NP samples): 120 positive, 320 negative. Sensitivity overall: 75.8% (91/120), sensitivity at Ct<25: 98.8% (86/87). Specificity: 100% (320).</p>	Nasopharyngeal	-	Unknown	06/05/2022
1988	BCOV-502	Inzek International Trading B.V.	Biozek covid-19 Antigen Rapidtest BCOV-502	<p>Prospective clinical field study</p> <p>Study in the Netherlands involving a local public health authority (n=950, PCR positive = 61), NP swab; sensitivity overall: 85.25%; specificity: 99.78%.</p>	Nasopharyngeal	-	Nucleocapsid protein	04/03/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				<p>Prospective clinical field study Study in the Netherlands among healthcare workers (n=294, PCR positive = 44), NP swab; sensitivity overall: 81.8%, sensitivity Ct<30: 91.9%; specificity: 99.7%.</p>				
1764	G10313	JOYSBIO (Tianjin) Biotechnology Co., Ltd. (manufacturer)	SARS-CoV-2 Antigen Rapid Test Kit (Colloidal Gold)	<p>Prospective clinical field study Study in Czechia, N=225 (90 RT-PCR positive), 60.3% symptomatic patients. Test parameters for a subgroup of symptomatic patients: sensitivity 92% (80.8–97.8), specificity 97.6% (91.5–99.7). Test parameters for a subgroup of asymptomatic patients: sensitivity 100% (54.1–100), specificity 100% (95.5–100). Source.</p> <p>Prospective clinical field study Study in Italy (nasal swab) including asymptomatic or mild symptomatic participants, compared against RT-PCR from NP swab. Sample size: 115 positive, 386 negative samples. Overall sensitivity: 98.3%, specificity 99.2%</p>	Nasal	-	Nucleocapsid protein	10/05/2021
1353	AGSWNSA21-01, AGSWNSA21-02, AGSWNSA21-04, AGSWNSA21-05	LINKCARE (NANTONG DIAGNOS BIO)	COVID-19 Antigen Test Kit (Colloidal Gold)	<p>Prospective clinical field study Prospective study in Spain, N = 504 nasal samples (385 negative and 115 positive), performed by University Hospital Son Espases. Sensitivity: 96.33% (CI95 0.91-0.99); specificity: 100%; compared against PCR Ct ≤ 30.</p> <p>Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.04%</i>.</p>	Nasal, Nasopharyngeal	-	Nucleocapsid protein	21/12/2021
1268	SPEC-32312 R7 ART-00571 R13	LumiraDX	LumiraDx SARS-CoV-2 Ag Test	<p>Prospective clinical field study Evaluation by SKUP - Scandinavian evaluation of laboratory equipment for point of care testing. Total sample size: 448; 83 positive samples and 365 negative samples. For nasal specimen: sensitivity of 87% (79-92) and specificity of 99.5% (98.3-99.9). For nasopharyngeal specimen: sensitivity of 90% (83-95) and specificity of 97.8% (96.0-98.8). Source.</p>	Nasal	-	Nucleocapsid protein	17/02/2021

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				<p>FIND prospective evaluation study Germany (8 Oct 2021): 761 samples, NP swab. Clinical sensitivities: Days < 7: 86.4%; Ct ≤ 33: 87.2%; Ct < 25: 92.6%; Clinical specificity: 99.3%. Source.</p> <p>Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.8%</i>.</p>				
2640	11811125	Mologic Ltd	COVIOS Ag COVID-19 Antigen Rapid Diagnostic Test	<p>FIND prospective evaluation study Germany: Symptomatic and asymptomatic (n=649, PCR positive = 191), nasal and nasal-mouth-throat swab; sensitivity overall: 90.6%, sensitivity Ct ≤ 25: 96.4%; specificity: 100%.</p>	Nasal	-	Nucleocapsid protein	08/12/2021
2241		NESAPOR EUROPA SL	MARESKIT COVID-19 ANTIGEN RAPID TEST KIT	<p>Prospective clinical field study Prospective study in Spain; Nasal test compared to nasal PCR. Sensitivity 95.24% (Ct<30), Specificity 100%.</p>	Nasal	-	Nucleocapsid protein	23/07/2021
1880	NGB-COV-S23-202 (French market), NGB-COV-S23-203 (German market)	NG Biotech	Ninonasal	<p>Prospective clinical field study Prospective study in France for NP and nasal swabs: NP sensitivity 89% (75/84), specificity 99% (92/93). Nasal sensitivity 98% (125/128), specificity 99% (388/390)</p>	Nasal, Nasopharyngeal	-	Nucleocapsid protein	10/11/2021
1593		OSANG Healthcare Co., Ltd.	GeneFinder COVID-19 Ag Rapid Test	<p>Prospective clinical field study Independent prospective study carried out in a public hospital in Italy. Sample size: 100 positive and 450 negative samples. Overall sensitivity: 96%, specificity: 99.8%.</p>	Nasopharyngeal	-	Nucleocapsid protein	22/07/2022
2741	O1-01SA-SN	OSANG Healthcare Co., Ltd.	GeneFinder COVID-19 Ag Plus Rapid Test	<p>Prospective clinical field study Independent prospective evaluation study carried out in Hospital Pugliese Ciaccio, Italy. Sample type: NP swab; sample size: 100 positive, 400 negative; sensitivity: 94%; specificity: 100%.</p> <p>Prospective clinical field study Independent prospective field study in Italy: 151 positive samples, 452 negative samples. Sensitivity: 96.03%; Specificity: 99.78%.</p>	Nasopharyngeal	Nasal, Oropharyngeal	Nucleocapsid protein	21/12/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1097		Quidel Corporation	Sofia SARS Antigen FIA	<p>Prospective clinical field study</p> <p>Validation study in France, nasopharyngeal swabs. Sensitivity 84.44% (76/90), specificity 99.19% (491/495).</p> <p>Prospective clinical field study</p> <p>Independent prospective clinical field study in the Netherlands among symptomatic (n=733, PCR positive 144); NP swab; sensitivity overall: 84.0%, sensitivity Ct ≤ 30: 90.1%, sensitivity Ct ≤ 25: 92.5%; specificity overall: 99.8%</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	17/02/2021
1604	9901-NCOV-01G	Roche (SD BIOSENSOR) (manufacturer)	SARS-CoV-2 Rapid Antigen Test	<p>Prospective clinical field study</p> <p>Independent prospective clinical field study in the Netherlands among symptomatic (n=970, PCR positive 186); NP swab; sensitivity overall: 84.9%, sensitivity Ct ≤ 30: 94.3%, sensitivity Ct ≤ 25: 99.1%; specificity overall: 99.5%. Source.</p>	Nasopharyngeal	-	Nucleocapsid protein	10/05/2021
2228	9901-NCOV-03G	Roche (SD BIOSENSOR) (manufacturer)	SARS-CoV-2 Rapid Antigen Test Nasal	<p>Prospective clinical field study</p> <p>Study by the Charité Berlin in Germany for nasal samples and reference RT-PCR with NP/OP samples. Study size: 150 RT-PCR positive and 546 RT-PCR negative samples. Overall Sensitivity: 82.7% and of 97.8% for Ct ≤ 24 (91 samples). Overall Specificity: 99.1%</p>	Nasal	-	Nucleocapsid protein	07/07/2021
2052	Q-NCOV-04G	SD BIOSENSOR Inc. (manufacturer)	STANDARD Q COVID-19 Ag Test Nasal	<p>Prospective clinical field study</p> <p>Independent prospective clinical field study in the Netherlands among symptomatic (n=628, PCR positive 118); NP swab; sensitivity overall: 78.0%, sensitivity Ct ≤ 30: 84.4%, sensitivity Ct ≤ 25: 90.3%; specificity overall: 99.6%.</p> <p>Retrospective in vitro study</p> <p>Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.52%</i>.</p>	Nasopharyngeal	-	Nucleocapsid protein	22/07/2022
344	F-NCOV-01G	SD BIOSENSOR Inc. (manufacturer)	STANDARD F COVID-19 Ag FIA	<p>Prospective clinical field study</p> <p>Independent prospective clinical field study in the Netherlands among symptomatic (n=628, PCR positive 118); NP swab; sensitivity overall: 78.0%, sensitivity Ct ≤ 30: 84.4%, sensitivity Ct ≤ 25: 90.3%; specificity overall: 99.6%.</p> <p>Retrospective in vitro study</p> <p>Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.52%</i>.</p>	Nasopharyngeal	-	Nucleocapsid protein	17/02/2021
345	Q-NCOV-01G	SD BIOSENSOR Inc. (manufacturer)	STANDARD Q COVID-19 Ag Test	<p>Prospective clinical field study</p> <p>Study in Portugal: 80 samples from symptomatic individuals (27 PCR positive and 53 negative by PCR). Sensitivity: 70% (95%IC50-86); specificity: 100% (95%IC 93-100). TCID50/ml</p>	Nasopharyngeal	-	Nucleocapsid protein	17/02/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				0,68x 102 and CT<25. FIND prospective evaluation study Germany (10 Dec 2020): 1263 samples, NP swab. Clinical sensitivities: Days ≤ 7: 80%; Ct ≤ 33: 87.8%; Ct ≤ 25: 100%; Clinical specificity: 99.3%. Source .				
1592		Shenzhen Lifotronic Technology Co., Ltd.	Antigen Rapid Test Ag SARS-CoV-2	Prospective clinical field study Independent prospective study carried out at a public hospital in Catanzaro, Italy. Sample size: 100 positive and 450 negative samples. Sensitivity: 93% (95% CI: 86.25% - 96.57%), specificity: 99.78% (95% CI: 98.75% - 99.96%).	Nasopharyngeal	-	Nucleocapsid protein	22/07/2022
2017	SC0201	Shenzhen Ultra-Diagnostics Biotech Co., Ltd. (manufacturer)	SARS-CoV-2 Antigen Test Kit	Prospective clinical field study Study in Slovenia: sensitivity in unselected symptomatic population: 86.4% (172 RAT pos. / 199 RT-PCR pos.), sensitivity of 97.8% at Ct ≤ 25. Specificity: 99.1% (1972 RAT neg. / 1990 RT-PCR neg.), NP swab.	Nasopharyngeal	Nasal ! Saliva	Nucleocapsid protein	10/05/2021
1466	2276, 2276/20	TODA PHARMA	TODA CORONADIAG Ag	Prospective clinical field study Study in France: NP swabs, sensitivity : 96.1-100%, specificity 99.2-100%. Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%</i> .	Nasopharyngeal	Nasal	Nucleocapsid protein	10/05/2021
2111	VCD 16 {...}	VivaChek Biotech (Hangzhou) Co., Ltd	SARS-CoV-2 Ag Rapid Test	Prospective clinical field study Study at General Hospital Jesenice in Slovenia; Nasal specimens. Total of 472 samples: 113 positive and 359 negative samples. Sensitivity: 85.84%, specificity: 99.72%.	Anterior nasal	-	Nucleocapsid	10/06/2022
1957	LCV03 {...}	Zhuhai Lituo Biotechnology Co., Ltd. (manufacturer)	COVID-19 Antigen Detection Kit (Colloidal Gold)	Prospective clinical field study Independent prospective field study at a public hospital in Slovenia; Nasal specimens; sensitivity 189/191 PCR positives: 98.95%, specificity 403/404: 100%. Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI)	Nasal, Nasopharyngeal	-	Nucleocapsid protein	14/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since <i>D/M/Y</i>
				in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 100%.				
2201	2061708 (1test/kit), 2061701 (20tests/kit),	Zybio Inc.	SARS-CoV-2 Antigen Assay Kit (Colloidal Gold Method)	Prospective clinical field study Independent prospective field study at a public hospital in Slovenia; nasal samples. Study population: unselected hospital patients, 107 positive and 417 negative samples (as defined by RT-PCR testing of matched NP swabs). Sensitivity: 88.8%; specificity: 99%.	Nasal	-	Nucleocapsid protein	04/03/2022

CATEGORY A.2: COVID-19 LABORATORY-BASED ANTIGENIC ASSAYS

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since <i>D/M/Y</i>
1960	311490 (reagent), 311491 (control), 311492 (buffer)	DIASORIN	LIAISON SARS-CoV-2 Ag assay	Prospective clinical field study Study in Belgium (n=414, PCR positive = 204, PCR negative = 210), NP swab. Sensitivity Ct ≤ 35: 73.4%, sensitivity Ct ≤ 25: 96.4%; specificity: 100%. Prospective clinical field study Study in Italy, symptomatic and asymptomatic (n=378, PCR positive = 46), NP swab. Overall sensitivity: 84.8%, sensitivity Ct ≤ 25: 100%; specificity: 99.4%. Prospective clinical field study Study in Italy (n=1075, PCR positive = 23), NP swab; sensitivity Ct ≤ 30: 90.5%; specificity: 99.8%. Prospective clinical field study Independent field study in the Netherlands (n=980, PCR positive n=98), NP+OP swab; sensitivity overall: 82.7%, sensitivity Ct ≤ 30: 91.9%; specificity overall: 99.1%.	Nasopharyngeal	Nasal	Nucleocapsid protein	20/10/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Evaluated specimen type(s) <i>Eligible for issuing EU Digital COVID certificates</i>	Other specimen type(s) ¹⁷ <i>Not evaluated</i>	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2124	260340	Fujirebio	Lumipulse G SARS-CoV-2 Ag	<p>Prospective clinical field study</p> <p>Study in Belgium, NP samples: 102 positive samples, 400 negative samples (including 100 hospitalized patients). Sensitivity: 93%, specificity: 99%.</p> <p>Prospective clinical field study</p> <p>Study in Italy; sample size (NP): 194 positive and 400 negative. Sensitivity (overall): 79.9% (155/194); sensitivity (Ct ≤ 25): 100% (87/87); specificity: 99.3% (397/400).</p>	Nasopharyngeal	-	Nucleocapsid protein	08/04/2022
1200	619 9941 (reagent pack), 619 9942 (calibrator)	Ortho Clinical Diagnostics	VITROS Immunodiagnostic Products SARS-CoV-2 Antigen	<p>Prospective clinical field study</p> <p>Study in Belgium: 80 positive NP samples (sensitivity 100%), 108 negative samples (specificity 100%).</p> <p>Prospective clinical field study</p> <p>Study in France: 107 positive NP samples with Ct ≤ 35 (sensitivity 93.5%), 1614 negative samples (specificity 100%).</p> <p>Retrospective in vitro study</p> <p>A retrospective study including 134 positive NP samples with Ct ≤ 35 (sensitivity 82.8%).</p>	Nasopharyngeal	Nasal	Nucleocapsid protein	10/02/2022
2156	09345299190	Roche Diagnostics GmbH	Elecsys® SARS-CoV-2 Antigen2156	<p>Prospective clinical field study</p> <p>Study in Germany: Total N: 3139 (2747 negative, 392 positive). Relative specificity overall 99.9%; relative sensitivity (n=390) overall 92.5% (Ct ≤ 26).</p>	Nasopharyngeal	Nasal, Oropharyngeal	Nucleocapsid protein	20/10/2021

Category B: COVID-19 antigen tests evaluated by retrospective in vitro studies

The clinical performance of the following antigen tests listed under “Category B” has been evaluated by retrospective in vitro studies, meeting the criteria and definitions as agreed by the Health Security Committee on 21 September 2021.

Devices highlighted in blue are identical in design and construction but are, for example, branded or distributed under a different name. The results of validation studies may be transferred between devices that are identical in design and construction.

Important notes to be taken into account by EU Member States:

- ➔ In case of retrospective in vitro evaluation studies carried out by the Paul-Ehrlich-Institut in Germany, only the sensitivity of the device has been evaluated. The specificity as reported by the manufacturer has been indicated in the corresponding column. EU Member States should pay particular attention to the issuance of EU Digital COVID recovery certificates based on the result of these devices, as the specificity of the device has thus not been evaluated by an independent validation study meeting the agreed criteria.
- ➔ In general, retrospective in vitro studies do not aim to evaluate the clinical performance of an antigen test based on a specific specimen type. Therefore, the clinical performance of devices listed under Category B *cannot* be linked to a specific specimen type, which should be taken into consideration by countries when using these antigen tests for the issuance of EU Digital COVID certificates. Instead, the table below makes a general reference to the specimen type(s) that can be used for the device as stated in the Instructions For Use of the device.

CATEGORY B.1: COVID-19 RAPID ANTIGEN TESTS

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2374	7427245282658	ABIOTEQ	Cora Gentest-19	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.8%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal, Throat	Nucleocapsid protein	20/10/2021
2579	ABT-IDT-B367	AccuBioTech Co.,Ltd	Accu-Tell SARS-CoV-2 Ag Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasopharyngeal	Nucleocapsid protein	20/10/2021
1865	L031-12515, L031-125D5	Acon Biotech (Hangzhou) Co., Ltd	Flowflex SARS-CoV-2 Antigen Rapid Test (Nasal/Saliva)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal ! Saliva	Nucleocapsid protein	10/02/2022
1468	L031-11815	ACON Laboratories, Inc. (manufacturer)	Flowflex SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 98.7%.</i>	Nasal	Nucleocapsid protein	10/05/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2130	1166-25	Affimedix Inc. (manufacturer)	TestNOW® - COVID-19 Antigen Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	10/05/2021
1304	RT2950 (1 cassette), RT2955 (5 cassettes), RT2951 (10 cassettes)	AMEDA Labordiagnostik GmbH (manufacturer)	AMP Rapid Test SARS-CoV-2 Ag	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	17/02/2021
1822	A6061204, A6061281 (1 cassette), A6061282 (5 cassettes), A6061283 (10 cassettes)	Anbio (Xiamen) Biotechnology Co., Ltd	Rapid COVID-19 Antigen-Test (colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Throat	Nucleocapsid protein	10/05/2021
1736		Anhui Deep Blue Medical Technology Co., Ltd (manufacturer)	COVID-19 (SARS-CoV-2) Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.8%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
1815			COVID-19 (SARS-CoV-2) Antigen Test Kit (Colloidal Gold) – Nasal swab		Nasal, Anterior nasal		10/05/2021
2089	FCB-103	Anhui Formaster Biosci Co., Ltd.	New Coronavirus (COVID-19) Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.5%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
1926		ARISTA Biotech Pte.LTD.	ARISTA™ COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal	Nucleocapsid protein	08/04/2022
1618	A03-50-422	Artron Laboratories Inc.	Artron COVID-19 Antigen Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	14/07/2021
1654	AM3474-K (20T), AM3476-K (25T)	Asan Pharmaceutical Co., Ltd	Asan Easy Test COVID-19 Ag	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.7%.</i>	Nasal	Unknown	10/05/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
770	COV-S23	Assure Tech. (Hangzhou) Co., Ltd.	ECOTEST COVID-19 Antigen Rapid Test Device	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; Manufacturer specificity of 99.2%.	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
2350	COV-S23		ECOTEST COVID-19 Antigen Rapid Test Device		Nasopharyngeal, Oropharyngeal		23/07/2021
1800	03760097080046	Avalun	Ksmart® SARS-COV2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; Manufacturer specificity of 99.32%.	Nasopharyngeal	Unknown	07/07/2021
2101	88 99 19	AXIOM Gesellschaft für Diagnostica und Biochemica mbH	COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 100%.	Nasal, Nasopharyngeal, Throat	Nucleocapsid protein	10/05/2021
2807	HGCG134S01 {...}	Beijing Hotgen Biotech Co., Ltd	Coronavirus (2019-nCoV)-Antigentest	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 98.88%.	Nasal	Nucleocapsid protein	21/01/2022
1870	AT120/20 (Model A/B)	Beijing Hotgen Biotech Co., Ltd	Novel Coronavirus 2019-nCoV Antigen Test (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 99.76%.	Nasal, Throat ! Saliva	Nucleocapsid protein	10/05/2021
2072	C3042	Beijing Jinwofu Bioengineering Technology Co.,Ltd.	Novel Coronavirus (SARS-CoV-2) Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 100%.	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal ! Saliva	Nucleocapsid protein	14/07/2021
1331	CG27 {...}	Beijing Lepu Medical Technology Co., Ltd (manufacturer)	SARS-CoV-2 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 99.26%.	Nasal, Nasopharyngeal	Nucleocapsid protein	17/02/2021
2494	00013C	Beijing O&D Biotech Co., Ltd.	COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 98.67%.	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
2858		Bioscience (Tianjin) Diagnostic Technology Co.,Ltd	Novel Coronavirus (2019-nCoV) Antigen Rapid Detection	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; Manufacturer specificity of 99.2%.	Nasal	Nucleocapsid protein	10/06/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2247	COVAGO {...}	BioGnost Ltd	CoviGnost AG Test Device 1x20	Retrospective in vitro study Study in Croatia: 300 NP samples, symptomatic (<7 dps): 200 PCR+ samples (range Ct 16-30), Ct ≤ 30: sensitivity 96.5%. 100 PCR- samples: specificity 100%.	Nasopharyngeal	Unknown	23/07/2021
2230	209.01.25.01	Biohit Healthcare (Hefei) Co., Ltd.	SARS-CoV-2 Antigen Rapid Test (Colloidal Gold Method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.49%.</i>	Nasal	Nucleocapsid protein	08/12/2021
1286	AT043/20 (Ref 34)	Biohit Healthcare (Hefei) Co., Ltd.	SARS-CoV-2 Antigen Rapid Test Kit (Fluorescence Immunochromato-graphy)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.9%.</i>	Anterior Nasal	Nucleocapsid protein	23/07/2021
1599	1509A-I	Biomerica Inc.	Biomerica COVID-19 Antigen Rapid Test (nasopharyngeal swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.7%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	07/07/2021
1242	RG1901DG	BIONOTE	NowCheck COVID-19 Ag Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.6%.</i>	Nasal, Nasopharyngeal	Unknown	07/07/2021
2067	400-678-8982	BIOTEKE CORPORATION (WUXI) CO., LTD	SARS-CoV-2 Antigen Test Kit (colloidal gold method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 99.28%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
2013	RTB25CoV2C	Biotal Health S.L.U. BIOTAL HEALTH S.L.U	biotal SARS-CoV-2 Ag Card	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 99.28%.</i>	Nasopharyngeal	Nucleocapsid protein	23/07/2021
1236	COV-19C25	BTNX Inc (manufacturer)	Rapid Response COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
2696	C2104N01	Cesna Biyoteknoloji Araştırma Geliştirme Laboratuvar Sist. İnş. Müh. Dan. San. Tic. Ltd. Şti.	CHECK UP SARS-COV-2 NASAL ANTIGEN RAPID TEST	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.8%.</i>	Nasal	Nucleocapsid protein	21/12/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2746	C2104N02	Cesna Biyoteknoloji Araştırma Geliştirme Laboratuvar Sist.İnş.Müh.Dan.San.Tic.Ltd.Şti.	CHECK UP SARS-COV-2 NASOPHARYNGEAL RAPID ANTIGEN TEST	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.7%.</i>	Nasopharyngeal	Nucleocapsid protein	21/12/2021
2588		Changzhou Biowin Pharmaceutical Co.,Ltd.	Novel Coronavirus(COVID-19) Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.3%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/06/2022
1691	CCOV-201	Chil Tıbbi Malzeme Sanayi ve Ticaret Limited Şirketi	CHIL COVID-19 Antigen Rapid Test (Nasopharyngeal / Oropharyngeal Swab-Casette)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.57%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
2150		Chongqing M&D Biotechnology Co. Ltd	2019-nCoV Antigen Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal	Nucleocapsid protein	20/10/2021
2449	ICOV-502	Citest Diagnostics Inc.	COVID-19 Antigen Rapid Test (Nasal Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasal	Nucleocapsid protein	10/06/2022
1581	R0182C	CTK Biotech, Inc (manufacturer)	OnSite COVID-19 Ag Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	07/07/2021
2242	CV19IC	DNA Diagnostic (manufacturer)	COVID-19 Antigen Detection Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.56%.</i>	Nasal	Nucleocapsid protein	23/07/2021
2756	CV19AG	DNA Diagnostic (manufacturer)	SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.3%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	21/01/2022
2273	3715270	Dräger Safety AG & Co. KGaA	Dräger Antigen Test SARS-CoV-2	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 99.6%.</i>	Nasal	Nucleocapsid protein	20/10/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2533	DNK-1425-1	Dynamiker Biotechnolgy(Tianjin) Co., Ltd. (manufacturer)	Dynamiker SARS-CoV-2 Ag Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.1%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
1243	EGCV0101 {...}	Edinburgh Genetics Limited (manufacturer)	Edinburgh Genetics ActivXpress+ COVID-19 Antigen Complete Testing Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.24%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
2724	PCSYHN02	Fosun Diagnostics (Shanghai) Co.,Ltd., China	Fosun Covid-19 Ag Card	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.7%.</i>	Nasopharyngeal	Nucleocapsid protein	04/03/2022
1739	EBS 1020	Eurobio Scientific (manufacturer)	EBS SARS-CoV-2 Ag Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.1%.</i>	Nasal	Nucleocapsid protein	07/07/2021
1855	3985	GA Generic Assays GmbH	GA CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasopharyngeal	Nucleocapsid protein	23/07/2021
1244	COVAG025	GenBody Inc	GenBody COVID-19 Ag Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94% at Ct ≤ 25; <i>Manufacturer specificity of 99.19%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	08/04/2022
2642	CoVSLFA-20	Genobio Pharmaceutical Co., Ltd.	Virusee® SARS-CoV-2 Antigen Rapid Test (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	08/12/2021
2012	52025 {...} 52026 {...} 52027 {...} 52104 {...} 52112 {...} 52129 {...}	Genrui Biotech Inc (manufacturer)	SARS-CoV-2 Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.02%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	07/07/2021
1253	P2004	GenSure Biotech Inc (manufacturer)	GenSure COVID-19 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal	Unknown	10/05/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2853	P2004s	GenSure Biotech Inc (manufacturer)	GenSure COVID-19 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal ! Saliva	Nucleocapsid protein	10/02/2022
2183	CG2061	Getein Biotech, Inc. (manufacturer)	One Step Test for SARS-CoV-2 Antigen (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.71%.</i>	Nasal ! Saliva	Nucleocapsid protein	16/06/2021
1820	CG20615	Getein Biotech, Inc. (manufacturer)	SARS-CoV-2 Antigen (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.71%.</i>	Nasal ! Saliva	Nucleocapsid protein	14/07/2021
2695	600008	Glallergen CO., LTD.	Novel Coronavirus (2019-nCoV) Antigen Test Kit (Colloidal gold immunochromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.02%.</i>	Nasal	Nucleocapsid protein	21/12/2021
1197	CG123005	Goldsite Diagnostic Inc.	SARS-CoV-2 Antigen Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal ! Other	Nucleocapsid protein	14/07/2021
1216	LS-C-T-009	Guangdong Longsee Biomedical Co., Ltd.	2019-nCoV Ag Rapid Detection Kit (Immuno-Chromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
1360	BE0040, BE0041	Guangdong Wesail Biotech Co. Ltd (manufacturer)	COVID-19 Ag Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	17/02/2021
1324	0555C2X {...}	Guangzhou Decheng Biotechnology CO., Ltd	V-CHEK, 2019-nCoV Ag Rapid Test Kit (Immuno-chromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal	Nucleocapsid protein	07/07/2021
2257	INCP-502-N	Hangzhou AllTest Biotech Co., Ltd	SARS-CoV-2 Antigen Rapid Test (Nasal Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasal	Nucleocapsid protein	04/03/2022
2325			GSD NovaGen SARS-CoV-2 Ag Rapid Test (Nasal Swab)				22/07/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1876	302281	Hangzhou Biotest Biotech Co., Ltd	COVID-19 Antigen Rapid Test Cassette (Nasal Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasal	Nucleocapsid protein	08/12/2021
1610		Hangzhou Clongene Biotech Co., Ltd. (manufacturer)	COVID-19 Antigen Rapid Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.4% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal	Nucleocapsid protein	07/07/2021
1363	ICOV5002-B025		Covid-19 Antigen Rapid Test Kit		Nasal, Nasopharyngeal		17/02/2021
1365			COVID-19/Influenza A+B Antigen Combo Rapid Test		Nasopharyngeal		10/05/2021
2629		Hangzhou DIAN Biotechnology Co., Ltd.	COVID-19 Antigen Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 98.4%.</i>	Nasal, Nasopharyngeal	Unknown	21/12/2021
2862	INC-502	Hangzhou Funworld Biotech Co., Ltd	SARS-CoV-2 Antigen Rapid Test Device	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98%.</i>	Nasal, Nasopharyngeal ! Saliva	Nucleocapsid protein	08/04/2022
2885	P211138	Hangzhou GENESIS Biodetection and Biocontrol CO.,LTD	KaiBiLi COVID-19 Antigen Pro	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	10/06/2022
2979		Hangzhou Jucheng Medical Products Co., Ltd	SARS-CoV-2 Ag Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Anterior nasal	Nucleocapsid protein	08/04/2022
1215	303035	Hangzhou Laihe Biotech Co.	LYHER Novel Coronavirus (COVID-19) Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.7%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2022
2139	COV-201	Hangzhou Lysun Biotechnology Co. Ltd (manufacturer)	COVID-19 Antigen Rapid Test Device (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal	Nucleocapsid protein	10/05/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1945	COVG-602	Hangzhou Sejoy Electronics & Instruments Co.Ltd (manufacturer)	SARS-CoV-2 Antigen Rapid Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal	Nucleocapsid protein	08/12/2021
1952	COVG-602	Hangzhou Sejoy Electronics & Instruments Co.Ltd (manufacturer)	SARS-CoV-2 Antigen Rapid Test Cassette (nasal, nasopharyngeal, oropharyngeal, saliva)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.1%.</i>	Nasal, Nasopharyngeal, Oropharyngeal ! Saliva	Nucleocapsid protein	10/06/2022
1392	9010115	Hangzhou Testsea Biotechnology Co., Ltd.	Covid-19 Antigen Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.4%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	10/05/2022
2942	IFC-SCoV2-AG	Hangzhou Zheda Dixun Biological Gene Engineering Co., Ltd.	SARS-CoV-2 Nucleocapsid (N) Antigen Rapid Test Cassette (Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	06/05/2021
1929	HYT-G01	Hoyotek Biomedical Co., Ltd.	Corona Virus (COVID-19) Antigen Rapid Test (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasopharyngeal, Oropharyngeal	Unknown	20/10/2021
1759		Hubei Jinjian Biology Co., Ltd	SARS-CoV-2 Antigen Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.3%.</i>	Nasopharyngeal	Nucleocapsid protein	23/07/2021
1801	BT1389	Innova Medical Group.Inc	Innova SARS-CoV-2 Antigen Rapid Qualitative Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Anterior nasal, Nasal	Nucleocapsid protein	20/10/2021
2278	IN4658I	Innovation Biotech(Beijing) Co.Ltd	Coronavirus (SARS-Cov-2) Antigen Rapid Test Cassette (Nasal swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal	Nucleocapsid protein	20/10/2021
2419	ITP16010-TC1, ITP16010-TC25	InTec PRODUCTS, INC. (manufacturer)	Rapid SARS-CoV-2 Antigen Test (nasopharyngeal specimen)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal	Nucleocapsid protein	20/10/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1783	ITP16010-TC {...}	InTec PRODUCTS, INC. (manufacturer)	Rapid SARS-CoV-2 Antigen Test (nasopharyngeal/nasal specimen)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	08/1040/2021
2107	SC30107W {...}	Jiangsu Biopertectus Technologies Co., Ltd. (manufacturer)	Novel Corona Virus (SARS-CoV-2) Ag Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.15%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	14/07/2021
1920		Jiangsu Diagnostics Biotechnology Co., Ltd	COVID-19 Antigen Rapid Test Cassette (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Oropharyngeal, Throat	Nucleocapsid protein	14/07/2021
1899		Jiangsu Konsung Bio-Medical Science and Technology Co	COVID-19 Antigen Rapid Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.34%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/02/2022
2006	1031-02	Jiangsu Medomics medical technology Co.,Ltd. (manufacturer)	SARS-CoV-2 antigen Test Kit (LFIA)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.51%.</i>	Anterior nasal, Nasopharyngeal, Throat	Nucleocapsid protein	07/07/2021
2586	R220T001B0C0	Jiangsu Mole Bioscience CO., LTD.	SARS-CoV-2 Antigen Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.17%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	08/12/2021
2144	CO-05	Jiangsu Well Biotech Co., Ltd.	COVID-19 Ag Rapid Test Device	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal	Nucleocapsid protein	20/10/2021
2963		Jiangxi Province JinHuan Medical Instrument Co., LTD.	DREHA Novel Coronavirus (SARS-CoV-2) Antigen Rapid Detection Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal	Nucleocapsid protein	08/04/2022
1333		Joinstar Biomedical Technology Co. Ltd (manufacturer)	COVID-19 Rapid Antigen Test (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.1%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	17/02/2021
2555		IEDAU INTERNATIONAL GMBH	Covid-19 Antigen Schnelltest (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.1%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	08/12/2021

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2038		Koch Biotechnology (Beijing) Co., Ltd	COVID-19 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.3%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	22/07/2022
1266	B63000, LX-401301	Labnovation Technologies Inc.	SARS-CoV-2 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
2866		Lifecosc Biotech Limited	COVID-19 Antigen Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	06/05/2022
2128	P230 {...}	Lumigenex (Suzhou) Co., Ltd	PocRoc® SARS-CoV-2 Antigen Rapid Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.16%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
1267	71110	LumiQuick Diagnostics Inc (manufacturer)	QuickProfile™ COVID-19 Antigen Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.8%.</i>	Nasopharyngeal	Unknown	10/05/2021
1180	T00023 (single box), T00019 (25 pack)	MEDsan GmbH	MEDsan SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.8%.</i>	Nasopharyngeal, Oropharyngeal	Unknown	17/02/2021
2029	CP01810011, CP02150011	Merlin Biomedical (Xiamen) Co., Ltd.	SARS-CoV-2 Antigen Rapid Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.99%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	16/06/2021
1775	3011035	MEXACARE GmbH	MEXACARE COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.1%.</i>	Nasal	Nucleocapsid protein	07/07/2021
1190	0230005SP	möLab	mö-screen Corona Antigen Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.99%.</i>	Nasopharyngeal	Unknown	10/05/2021

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1481	07AG6020B	MP Biomedicals	Rapid SARS-CoV-2 Antigen Test Card	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.03%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	17/02/2021
2260	MGJGEN	Multi-G bvba	Covid19Check-NAS	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal	Nucleocapsid protein	10/02/2022
2301	500200	Nanjing Liming Bio-Products Co., Ltd.	StrongStep® SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.26%.</i>	Anterior nasal, Nasopharyngeal, Oropharyngeal ! Saliva	Nucleocapsid protein	08/12/2021
2506		Nanjing Norman Biological Technology Co., Ltd.	Novel Coronavirus (2019-nCoV) Antigen Testing Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasopharyngeal	Nucleocapsid protein	10/11/2021
2164	RQ005 {...}	Nanjing Synthgene Medical Technology Co., Ltd.	SARS-COV-2 Nucleocapsid (N) Antigen Rapid Detection Kit (Colloidal gold method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasopharyngeal	Nucleocapsid protein	21/01/2022
2200	B66000	NanoRepro AG	NanoRepro SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 98.4%.</i>	Anterior nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
1573		Nantong Egens Biotechnology Co.,Ltd	COVID-19 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal	Nucleocapsid protein	10/02/2022
2678		NDFOS Co., Ltd.	ND COVID-19 Ag Test	Retrospective in vitro study Independent retrospective study carried out by the University of Geno, Italy. Sample size: 100 positive and 300 negative samples. Panel distribution meet criteria. Overall sensitivity: 87% (79.02-92.24), sensitivity of 100% (91.24-100) at Ct ≤ 25, specificity: 100%.	Nasopharyngeal	Nucleocapsid protein	22/07/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2608	SVRAG	Neo-nostics (Suzhou) Bioengineering Co., Ltd.	COVID 19 Antigen Test Kit (Colloidal Gold Method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.19%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/02/2022
1501	COVID-19-NG08	New Gene (Hangzhou) Bioengineering Co., Ltd. (manufacturer)	COVID-19 Antigen Detection Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Nasal, Nasopharyngeal, Oropharyngeal ! Saliva, Sputum	Nucleocapsid protein	16/06/2021
1762	MY28	Novatech Tıbbi Cihaz Ürünleri Sanayi ve Ticaret A.Ş.	SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	14/07/2021
1199	CMA-031	Oncosem Onkolojik Sistemler San. ve Tic. A.S.	CAT	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 98.04%.</i>	Nasal	Nucleocapsid protein	10/05/2021
2116	67311, 67321, 67331, 67341, 67351, 67361, 67371	PerGrande Bio Tech Development Co., Ltd.	SARS-CoV-2 Antigen Detection Kit (Colloidal Gold Immunochromatographic Assay)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.11%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
2672		Pierenkemper GmbH	(SARS-CoV-2) Antigen Rapid Test COVIDENT (SWAB) COVID-19	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal Throat	Nucleocapsid protein	04/03/2022
1271	PR-FC13	Precision Biosensor Inc.	Exdia COVI-19 Ag	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.3%.</i>	Nasopharyngeal	Unknown	17/02/2021
2685	200063-20P	PRIMA Lab SA	COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.3%.</i>	Nasopharyngeal	Nucleocapsid protein	08/04/2022
1495	V1320, V1340	Prognosis Biotech (manufacturer)	Rapid Test Ag 2019-nCov	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.58%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	07/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1341	H100G	Qingdao Hightop Biotech Co., Ltd (manufacturer)	SARS-CoV-2 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.75%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	17/02/2021
2754	H100C	Qingdao Hightop Biotech Co., Ltd	SARS-CoV-2/Flu A+B/RSV Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.75%.</i>	Nasopharyngeal	Nucleocapsid protein	21/12/2021
2290	311590	Rapid Pathogen Screening, Inc	LIAISON® Quick Detect Covid Ag Assay	Retrospective in vitro study Independent validation study, in Italy; 100 positive and 100 negative samples. Sensitivity: 92.7% with Ct<25; specificity: 100%.	Nasal, Nasopharyngeal	Nucleocapsid protein	23/07/2021
1489	COV Ag-6012	Safecare Biotech (Hangzhou) Co. Ltd (manufacturer)	COVID-19 Antigen Rapid Test Kit (Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.4%.</i>	Nasal	Nucleocapsid protein	17/02/2021
1490	FCO-6032a (multi-windows cassette)		Multi-Respiratory Virus Antigen Test Kit (Swab) (Influenza A+B/COVID-19)				10/05/2021
2097	S3109E	Sansure Biotech Inc	SARS-CoV-2 Rapid Antigen Test (Colloidal Gold Method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.1%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	21/12/2021
1201	34	ScheBo Biotech	ScheBo SARS CoV-2 Quick Antigen	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasopharyngeal, Oropharyngeal ! Serum	Nucleocapsid protein	16/06/2021
2763	36	ScheBo Biotech	ScheBo SARS CoV-2 Quick ANTIGEN (Colloidal Gold Method)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.49%.</i>	Nasal	Nucleocapsid protein	21/01/2022
1319	SCVC02	SGA Medikal (manufacturer)	V-Chek SARS-CoV-2 Ag Rapid Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasal	Nucleocapsid protein	10/05/2021
1357	SCVC02		V-Chek SARS-CoV-2 Rapid Ag Test (Colloidal gold)				07/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2152	CG201A	Shenzhen CAS-Envision Medical Technology Co., Ltd.	SARS-CoV-2-Antigen Rapid Detection Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.5%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	08/12/2021
2415	R0043, R0044, R0045, R0046	Shenzhen Dymind Biotechnology Co., Ltd	SARS-CoV-2 Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.37%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	20/10/2021
2812	HRK-66 {...}	Shenzhen Huaree Technology Co.,Ltd	SARS-CoV-2 Antigen Rapid Test Kit (Immunochromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal	Nucleocapsid protein	06/05/2022
2414	203-020	Shenzhen Huian Biosci Technology Co., Ltd.	SARS-CoV-2 Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.1%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	20/10/2021
2941	150129	Shenzhen Kingfocus Biomedical Engineering Co., Ltd.	COVID-19 Antigen Detection Kit (Quantum Dots-Based Immunofluorescence Chromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.28%.</i>	Nasal	Nucleocapsid protein	08/04/2022
1813	K602-20	Shenzhen Kisshealth Biotechnology Co., Ltd	SARS-CoV-2 Antigen Test Kit (GICA)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.2%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
2109	GF102B1	Shenzhen Lvshiyuan Biotechnology Co., Ltd.	Green Spring SARS-CoV-2 Antigen-Rapid test-Set	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal ! Saliva	Nucleocapsid protein	10/05/2021
1967	MF-68	Shenzhen Microprofit Biotech Co., Ltd	SARS-CoV-2 Antigen Test Kit (Colloidal Gold Chromatographic Immunoassay)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	07/07/2021
1178	MF-60	Shenzhen Microprofit Biotech Co., Ltd.	SARS-CoV-2 Spike Protein Test Kit (Colloidal Gold Chromatographic Immunoassay)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Spike protein	23/07/2021

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1228	MF-63	Shenzhen Microprofit Biotech Co., Ltd.	SARS-CoV-2 Spike Protein Test Kit (Fluorescence Immunoassay)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasopharyngeal	Nucleocapsid protein, Spike protein (S1)	08/12/2021
2026	RNS92048B	Shenzhen Reagent Technology Co.,Ltd.	SARS-CoV-2 antigen IVD kit SWAB	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.1%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
1769	LFA0401-25N	Shenzhen Watmind Medical Co., Ltd (manufacturer)	SARS-CoV-2 Ag Diagnostic Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.12%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
1768	LFB0401-25N	Shenzhen Watmind Medical Co., Ltd (manufacturer)	SARS-CoV-2 Ag Diagnostic Test Kit (Immuno-fluorescence)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.13%.</i>	Nasal	Nucleocapsid protein	07/07/2021
1347	G86255, G86254, G86247	Shenzhen YHLO Biotech Co., Ltd.	GLINE-2019-nCoV Ag	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 99.85%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	08/12/2021
1780	SP-SW 106	Spring Healthcare Services AG	SARS-Cov-2 Antigen Rapid Test Cassette (swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasal	Nucleocapsid protein	10/06/2022
1114	CAGT025E0	Sugentech, Inc. (manufacturer)	SGTi-flex COVID-19 Ag	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	10/05/2021
2297	COVID19AGVCG	SureScreen Diagnostics	SARS-CoV-2 Rapid Antigen Test Cassette	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal	Nucleocapsid protein	20/10/2021
1942		Surge Medical Inc.	COVID-19 Antigen Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct < 25; <i>Manufacturer specificity of 99%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	21/01/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
3015		Suzhou Soochow University Saier Immuno Biotech Co., Ltd.	InstantSure Covid-19 Ag CARD	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.52%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	06/05/2022
3093	04A024	TBG BIOTECHNOLOGY XIAMEN INC.	SARS-CoV-2 Antigen Rapid Test (Nasal Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 99.68%.</i>	Nasal	Nucleocapsid protein	10/06/2022
2074	C011906	Triplex International Biosciences(China) CO.,LTD.	SARS-CoV-2 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.91%.</i>	Nasal, Nasopharyngeal, Oropharyngeal ! Saliva	Nucleocapsid protein	16/06/2021
1465	C011906	Triplex International Biosciences(China) CO.,LTD.	SARS-CoV-2 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal	Nucleocapsid protein	14/07/2021
1689	TICV03	TÜRKLAB TIBBİ MALZEMELER SAN. ve TİC. A.Ş.	Covid-19 Ag Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 99.54%.</i>	Nasal	Nucleocapsid protein	21/01/2022
2584	ICV03		INFO Covid-19 Ag Test				21/12/2021
1751	RTC03		RAPIDAN TESTER Covid-19 Ag Test				21/01/2022
1722	TCV03		TOYO Covid-19 Ag Test				21/01/2022
1443	VSCD02	Vitrosens Biotechnology Co., Ltd (manufacturer)	RapidFor SARS-CoV-2 Rapid Ag Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.05%.</i>	Anterior nasal, Nasal, Nasopharyngeal, Oropharyngeal Throat	Nucleocapsid protein	10/05/2021
2100	VCD 16 {...}	VivaChek Biotech (Hangzhou) Co., Ltd, China	Verino Pro SARS CoV 2 Ag Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	21/12/2021
1276	COVA1	Willi Fox GmbH	Willi Fox COVID-19 Antigen rapid test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.9%.</i>	Nasopharyngeal, Oropharyngeal ! Other	Nucleocapsid protein	10/06/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
2098	W-Ag03-20	Wuhan EasyDiagnosis Biomedicine Co., Ltd.	COVID-19 (SARS-CoV-2) Antigen-Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.26%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
2742	FP-318	Wuhan HealthCare Biotechnology Co. Ltd.	SARS-CoV-2 Antigen Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 90% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Nasal, Nasopharyngeal	Nucleocapsid protein	04/03/2022
1773	SF24025	Wuhan Life Origin Biotech Joint Stock Co., Ltd.	SARS-CoV-2 Antigen Assay Kit (Immuno-chromatography)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.13%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	14/07/2021
2090	CoV2Ag-25	Wuhan UNscience Biotechnology Co., Ltd. (manufacturer)	SARS-CoV-2 Antigen Rapid Test Kit	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.57%.</i>	Nasal, Nasopharyngeal, Oropharyngeal, Mid-turbinates	Nucleocapsid protein	07/07/2021
2143	COV-S31	Wuxi Biohermes Bio & Medical Technology Co., Ltd.	SARS-CoV-2 Antigen Test Kit (Lateral Flow Assay)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.02%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	20/10/2021
1763	CG01Ag-25	Xiamen AmonMed Biotechnology Co., Ltd (manufacturer)	COVID-19 Antigen Rapid Test Kit (Colloidal Gold)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.55%.</i>	Nasal	Nucleocapsid protein	10/05/2021
1278	1N40C5	Xiamen Boson Biotech Co. Ltd	Rapid SARS-CoV-2 Antigen Test Card	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 99.03%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	17/02/2021
1296	3021902	Zhejiang Anji Saianfu Biotech Co., Ltd (manufacturer)	AndLucky COVID-19 Antigen Rapid Test	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 94.1% at Ct ≤ 25; <i>Manufacturer specificity of 99%.</i>	Nasal, Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	10/05/2021
1295	COVG10 {...}		reOpenTest COVID-19 Antigen Rapid Test		Nasal, Nasopharyngeal		10/05/2021
2271	SARS-CoV-2019 Ag (N) {...}	Pantest SA (manufacturer)	Pantest Coronavirus Ag (Nasopharyngeal Swab)		Nasopharyngeal		08/12/2021
2687		Zephyr Biomedicals - Tulip Diagnostics	PerkinElmer COVID 19 Antigen Test (NS, NP)	Retrospective in vitro study Independent retrospective study carried out by the University of Chieti-Pescara, Italy. NP	Nasal, Nasopharyngeal	Nucleocapsid protein ! Spike	22/07/2022

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
				samples, 155 positive samples with a Ct distribution in agreement with criteria. Overall sensitivity: 80.64%, sensitivity 100% at Ct ≤ 25, specificity: 99%.		glycoprotein	
2684		Zhejiang GENE SCIENCE Co., Ltd	Novel Coronavirus (COVID-19) Antigen Detection Kit (Swab)	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 100% at Ct ≤ 25; <i>Manufacturer specificity of 98.73%.</i>	Nasopharyngeal, Oropharyngeal	Nucleocapsid protein	08/12/2021
1902	RDC-802	Zhuhai Encode Medical Engineering Co.,Ltd	ENCODE SARS-COV-2 Antigen Rapid Test Device	Retrospective in vitro study Positive evaluation by Paul-Ehrlich-Institut (PEI) in Germany: Sensitivity of 95% at Ct ≤ 25; <i>Manufacturer specificity of 100%.</i>	Anterior nasal, Nasal, Throat	Nucleocapsid protein	20/10/2021

CATEGORY B.2: COVID-19 LABORATORY-BASED ANTIGENIC ASSAYS

Device ID # ¹⁵	REF number ¹⁶	Name of submitting company (and role) ¹⁷	Commercial name of the device ¹⁷	Clinical performance of the device <i>As evaluated by independent validation studies, meeting the agreed criteria</i>	Specimen type(s) ¹⁷	SARS-CoV-2 Target protein	Included in the EU common list since D/M/Y
1982	ABCVA-020, ABCVA-020W	Absology Co., Ltd.	Absoludy COVID-19 Ag	Retrospective in vitro study Study in Italy, residual NP samples already tested by RT-PCR. Sample size: 148 positive samples (Ct ≤ 17-25: n=52; Ct ≤ 25-30: n=63; Ct ≤ 30-36: n=33); 300 negative samples. Sensitivity (overall): 80.4%; Specificity: 100%.	Nasopharyngeal	Nucleocapsid protein	10/06/2022

ANNEX II: Checklists and further guidance for manufacturers

CHECKLIST I

FOR MANUFACTURERS WISHING TO SUBMIT AN APPLICATION FOR A NEW DEVICE TO BE INCLUDED IN THE EU COMMON LIST OF COVID-19 ANTIGEN TESTS

For further details about the criteria and definitions referred to below, please see chapter 2.1

1. Is the device an **antibody test**?
2. Is the device a **antigen self-test**?
3. Is the device a **pooled antigen test**?
4. Is the device solely based on **sampling materials other than nasal, oropharyngeal or nasopharyngeal specimens** (e.g. is it a device that can only be used for saliva sampling)?

YES

*If the answer to **any of these questions** is **YES**, this means that the device is not eligible to be included in the EU common list.*

Your application will be rejected by the HSC.

NO

5. Does the device carry **CE-marking**?
6. Is the device **in use** in at least one of the 27 EU Member States?
7. Has the clinical performance of the device, based on nasal, oropharyngeal and/or nasopharyngeal swabs, been evaluated in at least one of the 27 EU Member States through an **independent validation study**?

NO

*If the answer to **any of these questions** is **NO**, this means that the device is not eligible to be included in the EU common list.*

Your application will be rejected by the HSC.

YES

*If the answer to **all of these questions** is **YES**, please continue with CHECKLIST II.*

CHECKLIST II

INDEPENDENT VALIDATION STUDIES CARRIED OUT IN A EU MEMBER STATE

1. Has the validation study be performed by an **independent laboratory**? Do you have a signed **declaration of conflict of interest**?

This is a laboratory not owned nor operated by the manufacturer or sponsor of the test, and which is not related to the operator by ownership, familial relationships, nor contractual or other relationships that result in the laboratory being controlled by or being under the common control of the operator.

*As part of your application, you should provide a **declaration of conflict of interest**, signed by the laboratory involved, containing details whether funding from a private partner or the manufacturer was received and stating that no one has personal interest in the manufacturer company, stock papers, or have worked there within the previous 3 years.*

2. Has the validation study be performed **objectively** and in the **public interest**?
3. Was a **public body** involved in the validation study? Do you have a **written endorsement** of this body?

A public body is a public entity (government institution, regional government office, public hospital) that is primarily state-funded and plays a role in public health.

*Ideally, the public body should be involved in the design, oversight and analysis of the study. However, as arrangements vary in Member States, you may also provide a **written endorsement** as part of your application, signed by a public body in case the study was performed by other organisations. Note that written endorsement by individuals are not accepted.*

NO

If the answer to **any of these questions** is **NO**, this means that the device is not eligible to be included in the EU common list.

Your application will be rejected by the HSC.

YES

4. Is the validation study based on a **prospective clinical field study design**, testing *unselected* symptomatic and asymptomatic participants for SARS-CoV-2 infection?

"Unselected" means no prior knowledge of SARS-CoV-2 diagnosis (e.g. determined by PCR); inclusion is allowed based on general possible COVID-like symptoms (or close contact with COVID-19 cases); and exclusion is allowed of children (e.g. <16 years) or for medical ethical permission reasons.

YES

Continue with CHECKLIST III.I

NO

5. Is the validation study based on a **retrospective in vitro study design**, testing the clinical performance of the device by using reference panels?

YES

Continue with CHECKLIST III.II

CHECKLIST III.I - PROSPECTIVE CLINICAL FIELD STUDIES

1. Does the study show a **sensitivity** of:
 - a. $\geq 80\%$ when testing unselected symptomatic participants within the first seven days after symptom onset or asymptomatic participants, where the diagnosis is confirmed by RT-PCR in independent field studies; or
 - b. $\geq 90\%$ for subjects with a $Ct \leq 25$, in independent evaluations of unselected participants?
2. Does the study show a **specificity** of $\geq 98\%$?
3. Has the **study population** been clearly defined in the study and application documents, stating the inclusion criteria of participants (symptomatic individuals, close contacts or asymptomatic individuals without known exposure)?

Recommendations: Ideally, the study population would include a minimum of 30 participants with a Ct value ≤ 25 . The distribution and the individual results of the Ct values of all positive PCR samples and the corresponding antigen test results must be provided. Positive samples should be taken within the first seven days after symptom onset or time of infection, if known, taking into account the incubation time.
4. Has the **study design** been clearly described (incl. RT-PCR protocol and the distribution of Ct values)? Samples should represent naturally occurring viral loads. Ideally, the sensitivity for stratified Ct values should be discernible from the report.

Recommendations: An analysis of the correlation of PCR positive/antigen positive samples should be provided, stratified by Ct value or IU/mL. A correlation of antigen negative/PCR negative samples and results (Ct values or IU/mL) of antigen negative/PCR positive samples should be included. The PCR protocol, including the type of platform, the gene targets amplified, and any reference materials used should be described. Sampling should be matched for antigen and NAAT testing, e.g., two simultaneous samples from each individual or optimally NAAT- and antigen testing from the same sample (e.g. from the eluate of one swab); the buffer/transport medium should be compatible for both NAAT and antigen testing; any volume change in the buffer/medium for sample uptake different from that of the proprietary assay, and/or between antigen and NAAT test should be clearly communicated. Each specimen type should be evaluated separately. All claimed specimen types should be compared with paired NAAT results from nasopharyngeal or oropharyngeal specimens. The analysis and the sample storage should be clearly described and carried out in line with the IFU. Test and reference samples should preferably be taken sequentially.
5. Is the study based on a target population of at least **100 fresh RT-PCR positive samples**, and at least **300 fresh RT-PCR negative samples**? Note that this number of samples should be reached for each specimen type evaluated.

In case of multiple smaller prospective clinical field studies that do not meet the minimum number of positive and/or negative samples separately but that do meet all the other criteria, the number of samples may be combined, provided that the different studies applied the same/similar methodologies and that sufficient details are provided on their study design.
6. Has **ethical approval** been provided by an institutional review board?



If the answer to **any of these questions** is **NO**, this means that the device is not eligible to be included in the EU common list.

Your application will be rejected by the HSC.

YES

If the answer to **all of these questions** is **YES**, the device may be eligible to be included in the EU common list.

CHECKLIST III.II - RETROSPECTIVE IN VITRO STUDIES

1. Does the study show a **sensitivity** of:
 - c. $\geq 80\%$ when testing all specimen in the reference panel are accepted; or
 - d. $\geq 90\%$ for subjects with a $Ct \leq 25$?
2. Does the study show a **specificity** of $\geq 98\%$ (as measured through the retrospective in vitro evaluation study or as specified by the manufacturer in the IFU)?
3. Is the **composition of the reference panel** as follows?
 - A panel of **at least 50 (pooled) clinical specimens** that cover naturally occurring viral loads with SARS-CoV-2 concentration ranging from approximately 1.1×10^9 to 4.2×10^2 genome copies per mL of specimen and Ct values between 17 and 36.
 - The whole evaluation panel has been subdivided into three subgroups: panel members, which are characterized by:
 - **Very high viral load** (Ct value 17-25; **35% (+/- 2%)** of the total number of pooled clinical specimens);
 - **High viral load** (Ct value 25-30; **45% (+/- 2%)** of the total number of pooled clinical specimens); and
 - **Moderate viral load** (Ct value 30-36; **35% (+/- 2%)** of the total number of pooled clinical specimens).
 - For each pool, up to ten clinical respiratory specimens (nasopharyngeal/oropharyngeal) obtained for routine diagnostics with different virus loads may be used. The sample volume per panel member should be sufficient to allow comparative evaluation with different tests included in the evaluation.
 - RT-PCR needs to be applied to determine the RNA load per pool.
 - For each antigen test and panel member, a pre-defined aliquot needs to be completely absorbed using the specimen collection device, e.g. swab, provided with the respective test.
 - Further steps needs to be strictly performed following the respective instructions for use (IFU).
 - The stability of the panel (antigen) must be considered throughout the preparation of the panel and the workflow up to the test.
4. Has **ethical approval** been provided by an institutional review board?



*If the answer to **any of these questions** is **NO**, this means that the device is not eligible to be included in the EU common list.*

Your application will be rejected by the HSC.

YES

*If the answer to **all of these questions** is **YES**, the device may be eligible to be included in the EU common list.*