

Diagnostic Solutions for COVID-19



- NowCheck COVID-19 Ag Test
- NowCheck COVID-19 IgM/IgG Test



What is COVID-19?

Coronavirus disease 2019 (COVID-19) is a new disease caused by a novel coronavirus (SARS-CoV-2). It is a respiratory illness that can spread from person to person. It is now rapidly spreading worldwide and has been announced as a pandemic by WHO. Presentations of COVID-19 have ranged from asymptomatic/mild symptoms to severe illness and mortality. The rapid diagnosis of COVID-19 is critical for the prevention and control of this pandemic.



Why Do We Need Antigen and Antibody Tests?



NowCheck COVID-19 Ag Test

The NowCheck COVID-19 Ag Test is a rapid chromatographic immunoassay for the qualitative detection of specific antigens to SARS-CoV-2 present in human nasopharynx or nasal cavity.

Specifications

- Specimen : Nasopharyngeal swab, Nasal swab
- Testing time : 15~30 min.
- Packing Unit : 25 Tests/Kit
- Storage Condition : 2-30°C (36-86°F)

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NowCheck

COVID-19 Ag Test

Test Procedure

Nasopharyngeal swab

- Insert a nasopharyngeal swab into the nostril of the patient, and rotate the swab over the posterior nasopharynx surface 3~4 times.
- Insert the swab into an extraction buffer tube, and stir it more than 5 times while squeezing the tube.



Remove the swab while squeezing the sides of the tube, and press the nozzle cap tightly onto the tube.



Test Result

of the nostrils, and rotate the swab against the nasal wall 5-6 times. Repeat in the other nostril, using the same swab.

Apply 3 drops of the extracted specimen

to the specimen hole of the test device.

or

Analysis

Nasal swab

Repeat in the other nostril, using the same swab.Insert the swab into an extraction buffer tube, and stir it more than 10 times while squeezing the tube.

After getting a patient to blow one's nose, insert a nasal swab into one

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Read the test result after 15~30 min.
The test can be read up to 30 min.



** Do not read test results after 30 min. It may give false results.



NowCheck COVID-19 Ag Test

Performance

:: Clinical evaluation



* Prospective diagnostic evaluation study in Brazil

A total of 400 nasopharyngeal swab specimens were collected from symptomatic patients. Results of the test were compared to the routine, diagnostic RT-PCR (Lab-developed assay based on the US CDC protocol) result.

Specimens from symptomatic patients (N=400)		Real-time PCR			
		Positive	Negative	Total	
NowCheck COVID-19 Ag Test	Positive	91	8	99	
	Negative	11	290	301	
	Total	102	298	400	
(linical Sancitivity - 20 20% (050% CT 21 70% 02 00%)					

ical Sensitivity = 89.2% (95%CI, 81.7% - 93.9%)

92.2% (95%CI, 84.8% - 96.2%) (Days ≤ 7 from symptom onset) 91.4% (95%CI, 83.9%% - 95.6%) (Ct ≤ 33)

Clinical Specificity = 97.3% (95%CI, 94.8% - 98.6%)

Reference : https://www.finddx.org/covid-19/sarscov2-eval-antigen/

* Prospective diagnostic evaluation study in Russia

A total of 50 nasopharyngeal swab specimens were collected and the results were compared to the diagnostic RT-PCR result, which is used for clinical management.

Nasopharyngeal swab specimens (N=50)		Real-time PCR		
		Positive	Negative	Total
NowCheck COVID-19 Ag Test	Positive	24	0	24
	Negative	1	25	26
	Total	25	25	50
	Clinical Sensitivity =	= 96% (95%CI, 79.65	5% - 99.9%)	

Clinical Specificity = 100% (95%CI, 86.28% - 100%)

:: Limit of Detection (LoD)

SARS-CoV-2 strain tested	Virus stock titer	Specimen type	LoD (Spiking titer)	Final working titer
NCCP 43326/	1 X 10 ^{6.2} TCID ₅₀ /ml	Direct nasopharyngeal swab	3.12 X 10 ^{2.2} TCID ₅₀ /ml	6.24 X 10 ^{1.2} TCID ₅₀ /ml
2020 /Korea		Direct nasal swab	1.56 X 10 ^{2.2} TCID ₅₀ /ml	5.37 X 10 ^{1.2} TCID ₅₀ /ml



NowCheck COVID-19 Ag Test

Performance

:: No Cross-reactivity

There was no cross-reactivity with potential cross-reactive substances except SARS-coronavirus.

Category	Sort	Result	Category	Sort	Result
Virus	SARS-coronavirus	POS		Haemophilus influenzae	NEG
	MERS-coronavirus	NEG	- 	Mycoplasma pneumoniae	NEG
	Human Coronavirus	NEG		Streptococcus pneumonia	NEG
	Influenza A	NEG	-	Streptococcus pyrogens	NEG
	Influenza B	NEG	-	Streptococcus salivarius	NEG
	Respiratory syncytial virus	NEG	Bacteria	Candida albicans	NEG
	Human Metapneumovirus	NEG	- -	Bordetella pertussis	NEG
	(hMPV)	NLU	-	Moraxella catarrhalis	NEG
	Parainfluenza virus	NEG NEG		Pseudomonas aeruginosa	NEG
	Rhinovirus			Staphylococcus epidermidis	NEG
-	Enterovirus	NEG		Mycobacterium tuberculosis	NEG
	Adenovirus	NEG		Pooled human nasal wash –	
	Human immunodeficiency virus lysate	NEG	Others	representative of normal respiratory microbial flora	NEG

:: Not Affected by Interfering Substances

There was no interference with endogenous/exogenous interfering substances.

Antibiotics

Mupirocin, Tobramycin, Erythromycin, Ciprofloxacin

Anti-inflammatory drugs

Acetaminophen, Acetylsalicylic acid, Ibuprofen

Autoimmune disease

Human anti-mouse antibody, Rheumatoid factor

Relevant medicines

Zanamivir, Oseltamivir, Artemetherlumefantrine, Doxycycline hyclate, etc.

Nasal sprays or drops

Neo-Synephrine, Afrin Nasal Spray,

Rhinocort, etc.

Serum protein

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Whole Blood, EDTA anticoagulated

NowCheck COVID-19 IgM/IgG Test

NowCheck COVID-19 IgM/IgG Test is a rapid chromatographic immunoassay for the qualitative detection of specific antibodies to SARS-CoV-2 present in human serum, plasma, or whole blood.

Specifications

- Specimen : Whole blood, Serum, Plasma
- Testing time : 10~15 min.
- Packing Unit : 25 Tests/Kit
- Storage Condition : 2-30°C (36-86°F)



[Using Capillary whole blood] • Collect the 20 µl of capillary 2 Add the collected blood 6 Add 3 drops (90 µl) of 4 Read the test result in whole blood to the black line to the specimen hole buffer vertically 10~15 min. 3 **20** µl 20 µl drops T min. C G M [Using serum/plasma/whole blood] 2 Add the collected blood 4 Read the test result in Collect the serum, plasma Add 3 drops (90 µl) of (10 µl) or whole blood (20 µl) to the specimen hole buffer vertically 10~15 min. with a micropipette Whole blood 20 µI Whole blood 20 µl Serum/plasma 3 **10** µl drops Serum/plasma min. **10** µl iowCheck OVID-19 IgM/IgG **Test Result** Negative Positive Invalid COVID-19 IgM/IgG COVID-19 IgM/IgG COVID-19 COVID-19 COVID-19 COVID-19 COVID-19 COVID-19 COVID-19 COVID-19 COVID-19 IgM/IgG IgM/IgG IgM/IgG IgM/IgG IgM/IgG IgM/IgG IgM/IgG IgM/IgG IgM/IgG Control Line 🕨 A A . • IgG Line 🕨 IgM Line ►

Test Procedure

NowCheck COVID-19 IgM/IgG Test

Performance

:: Clinical evaluation

- Diagnostic Sensitivity

A total of 105 serum specimens from patients confirmed by Real-time PCR were analyzed. The results are as shown below:

Positive specimens (N=31)		NowCheck COVID-19 IgM/IgG Test			
		IgM(+)	IgG(+)	IgM(+) and/or IgG	
	0 - 7	100% (1/1)	100% (1/1)	100% (1/1)	
Days (post-symptom onset)	8 - 14	66.7% (2/3)	100% (3/3)	100% (3/3)	96.8% (30/31)
	≥ 15	88.9% (24/27)	96.3% (26/27)	96.3% (26/27)	

Sensitivity 96.8% (30/31)

- Diagnostic Specificity

Negativo sposimons (N=74)	NowCheck COVID-19 IgM/IgG Test			
Negative specifiens (N=74)	IgM(-)	IgG(-)	IgM(-) and IgG(-)	
Confirmed by Real time-PCR	98.6% (73/74)	100% (74/74)	98.6% (73/74)	

The NowCheck COVID-19 IgM/IgG Test showed 96.8% of sensitivity and 98.6% of specificity, compared to the Real-time PCR method.

* Conducted in July 2020 at Department of Laboratory medicine, Seoul National University Bundang Hospital

** Reference: PowerChek[™] 2019-nCoV Real-time PCR kit (KCDC approved, FDA EUA)

:: Limit of Detection (LoD)

The LoD of NowCheck COVID-19 IgM/IgG Test for SN titer was 1:40.



Diagnostic Solutions for COVID-19

Key Features

ACCURATE	ANYWHERE	EASE OF USE
Good correlation with PCR assays	No specific equipment needed	Minimal operator -dependent steps
	MADKET DDECENICE	
FAST RESULTS	AND ACCEPTANCE	STABLE

Diagnostic Algorithm

For individuals suspected to have COVID-19



Ordering Information

Product No.	Product Name	Product Type	Packing Unit
RG1901DG (Nasopharyngeal) RG1901DGN (Nasal)	NowCheck COVID-19 Ag Test	Device	25 Tests/Kit
RB2901DG	NowCheck COVID-19 IgM/IgG Test	Device	25 Tests/Kit
RG1901CD	NowCheck COVID-19 Ag Control	Control	10 Tests/Kit
RB2901CD	NowCheck COVID-19 IgM/IgG Control	Control	10 Tests/Kit

