

Clinical Performance

1 Sensitivity and Specificity

The performance of Medomics COVID-19 Antigen Test Kit (LFIA) was established with 340 nasopharyngeal swabs collected from clinical patients who were suspected of COVID-19. Two nasopharyngeal swabs were collected from patients and one swab was tested directly using Medomics COVID-19 Ag Test Card. The real-time Polymerase Chain Reaction (RT-PCR) assay for the detection of SARS-CoV-2 was used as the comparator method to confirm the status of samples for this study.

The results are shown in the following table.

Medomics COVID-19 Ag test	RT-PCR		Total
	Positive	Negative	
Positive	115	2	117
Negative	4	219	223
Total	119	221	340

Sensitivity: 96.64% (91.62%-99.08%)
Specificity: 99.10% (96.77%-99.89%)
PPV: 98.29% (93.96%-99.79%)
NPV: 98.21% (95.47%-99.51%)
Accuracy: 98.24% (96.20%-99.35%)

Explanation of terms

Sensitivity = True Positives / (True Positives + False Negatives)

Specificity = True Negatives / (True Negatives + False Positives)

PPV (Positive Predictive Value) = True Positives / (True Positives + False Positives)

NPV (Negative Predictive Value) = True Negatives / (True Negatives + False Negatives)

Accuracy = (True Negatives + True Positives) / Total Samples

2 Limit of Detection (LoD) Study Results

Medomics COVID-19 Ag Test Card LoD was determined by evaluating different concentrations of inactivated SARS-CoV-2 virus. LoD studies determined the lowest detectable concentration of SARS-CoV-2 at which 100% of all (true positive) replicates test positive. The results are shown in the following table.

Concentration TCID ₅₀ /mL	Number Positive/Total	% Detected
10	20/20	100%

The Medomics COVID-19 Ag Test Card LoD was confirmed as 10 TCID₅₀/mL.

3 High Dose Hook Effect

No high dose hook effect on test performance was observed up to a concentration of 10⁵ TCID₅₀/mL of inactivated SARS-CoV-2 virus with the Medomics COVID-19 Ag Test Card.

4 Cross Reactivity (Analytical Specificity) and Microbial Interference

Cross reactivity and potential interference of Medomics COVID-19 Ag Test Card was evaluated by testing commensal and pathogenic microorganisms listed in the following table that may be present in the clinical samples. Each of the bacterium, viruses, and yeast were tested in triplicate in the absence or presence of inactivated SARS-CoV-2 virus.

Potential Cross-Reactant	Concentration Tested	Cross-Reactivity (Yes/No)
Influenza A H1N1	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Influenza A H3N2	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Influenza A H5N1	1.0 x 10 ⁵ TCID ₅₀ /mL	No

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Influenza A H7N9	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Influenza B Victoria	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Influenza B Yamagata	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Parainfluenza virus Type 1	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Respiratory syncytial virus	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Enterovirus CA16e	1.0 x 10 ⁵ TCID ₅₀ /mL	No
Adenovirus	1.0 x 10 ⁵ TCID ₅₀ /mL	No
<i>Mycoplasma pneumoniae</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Staphylococcus aureus</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Staphylococcus epidermidis</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Bordetella pertussis</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Legionella pneumophila</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Streptococcus pneumoniae</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Haemophilus influenzae</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Streptococcus pneumoniae</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Mycobacterium tuberculosis</i>	1.0 x 10 ⁶ CFU/mL	No
<i>Candida albicans</i>	1.0 x 10 ⁶ CFU/mL	No

5 Endogenous Interfering Substances Effect

A study was performed to evaluate and demonstrate that the endogenous substances naturally present or drugs that may be artificially introduced into clinical samples do not interfere with the detection of SARS-CoV-2 in the Medomics COVID-19 Ag Test Card at the concentrations listed below.

Substance	Potential Interfering Substances	Concentration	Interference (Yes/No)
Endogenous	Mucin	2 % w/v	No
	Whole Blood	5 % v/v	No
Nasal spray	Phenylephrine	0.05 mg/ml	No

or drops			
Nasal steroids	Dexamethasone	0.8 mg/ml	No
	Triamcinolone acetonide	0.8 mg/ml	No
	Budesonide	0.5 mg/ml	No