

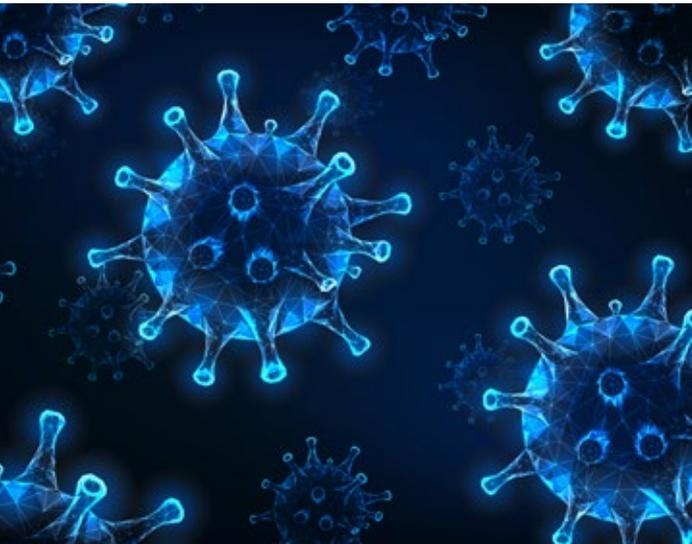
IVD



## **mariPOC<sup>®</sup>**

- Point-of-Care Testing for Respiratory Diagnostics

# mariPOC<sup>®</sup> - A fully automated continuous-feed immunoassay analyzer



The mariPOC<sup>®</sup> System is a compact solution for high-throughput, rapid Point-Of-Care (POC) testing needs that analyses swab samples fully automated. With a workflow of less than 3 minutes hands-on time and no need for additional instrumentation, it is an easy-to-use tool to test for pathogens anywhere by suitably trained individuals and in accordance with local regulations. It utilizes modern two-photon excitation fluorometry in combination with immunoassay-based testing to achieve a specificity and sensitivity so far only achieved by state-of-the-art immunoassay laboratories.

## Access to SARS-CoV-2 testing everywhere

The mariPOC<sup>®</sup> SARS-CoV-2 test supports detection of acute and contagious COVID-19 disease at the site of sampling. An indicative positive result from nasopharyngeal samples can be obtained after 20 minutes for highly positive infections, a final confirmative positive or negative result can be read after 90 minutes. Minimal work steps, automated analysis and the easy-to-use interface allow the use even with a large number of patients of up to 300 samples per 24 hours.

## Prepared for POC testing in winter

The mariPOC<sup>®</sup> Respi+ test allows point-of-care testing for a broad panel of respiratory-related pathogens covering SARS-CoV-2, Influenza, RSV, Adenovirus, Metapneumovirus, Parainfluenza, Coronavirus OC43 and *Streptococcus pneumoniae*. From just one

sample you get automated analysis for all these pathogens from nasopharyngeal swabs. The indicative positive result is available after 20 minutes for highly positive infections and a final confirmative result to identify if any of the pathogens is positive can be read after 2 hours.

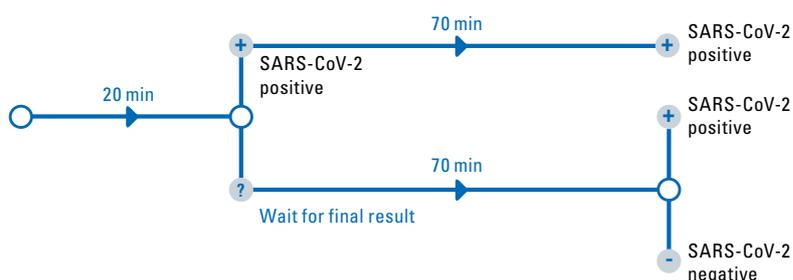


# Performance and result interpretation

## mariPOC® SARS-CoV-2

Pathogen	Sensitivity (N)	Specificity (N)
SARS coronavirus 2	92.3% (12/13) <sup>1</sup>	100% (198/198) <sup>1</sup>

<sup>1</sup> Compared to PCR



## mariPOC® Respi+

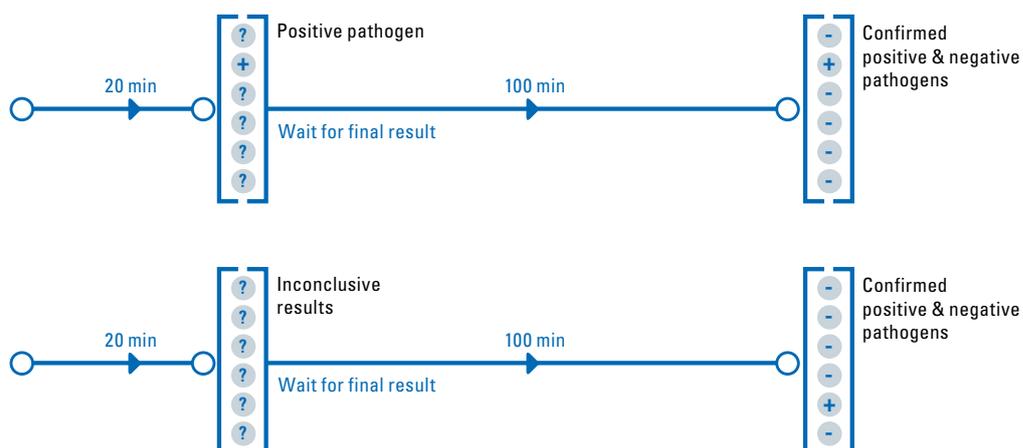
Pathogen	Sensitivity (N)	Specificity (N)
SARS coronavirus 2	92.3% (12/13) <sup>1</sup>	100% (198/198) <sup>1</sup>
Influenza A virus	92.3% (24/26) <sup>1</sup>	99.8% (870/872) <sup>1</sup>
Influenza B virus	87.5% (35/40) <sup>1</sup>	100% (152/152) <sup>1</sup>
Respiratory Syncytial Virus	88.6% (31/35) <sup>1</sup>	100% (123/123) <sup>1</sup>
Human metapneumovirus	77.8% (7/9) <sup>1</sup>	100% (195/195) <sup>1</sup>
Humona coronavirus OC43	NA	100% (159/160) <sup>1</sup>
<i>S. pneumoniae</i>	Similar <sup>2</sup>	No cross-reactions
Parainfluenza 1 virus	Similar <sup>3*</sup>	99% (198/200) <sup>1</sup>
Parainfluenza 2 virus	Similar <sup>3*</sup>	99% (198/200) <sup>1</sup>
Parainfluenza 3 virus	Similar <sup>3*</sup>	99% (200/202) <sup>1</sup>
Adenovirus	92.3% (24/26) <sup>3</sup>	100% (202/202) <sup>1</sup>

<sup>1</sup> Compared to PCR

<sup>2</sup> Compared to rapid antigen test

<sup>3</sup> Compared to time-resolved fluoroimmunoassay (TR-FIA)

\* Defined by antigen dilution series with TR-FIA due to low incidence of positive clinical samples



**Order Information:**

**Part No. 1204S | mariPOC® SARS-CoV-2**

Tests/plate: 308

Sample capacity: 308 samples/24 h

**Part No. 1184M | mariPOC® Respi+**

Tests/plate: 22

Sample capacity: 100 samples/24 h

**Part No. H-AD7-020 | mariPOC® AD7-020 analyzer**

System for use with mariPOC® tests

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In countries, where the IVD-CE mark is not applicable, mariPOC® is subject to local regulatory approval.  
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