

EC-Declaration of Conformity

According to Directive 98/79/EC on in-vitro-diagnostic devices, Annex III

product number: 243001N-10
product name: NADAL® COVID-19 IgG/IgM Test
classification: Other Products
manufacturer: nal von minden GmbH
Carl-Zeiss-Str. 12
47445 Moers

We herewith declare on our sole responsibility that all batches of above In-vitro-diagnostic device is conform with the Essential Requirements Annex I of the directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices. The product is suitable for the intended application (only professional users). Relevant standards and guidelines are applied.

This document is valid until 2022-05-25.

Moers, 05.03.2020



Sandra von Minden
CEO
nal von minden GmbH

nal von minden GmbH

Carl-Zeiss-Str. 12
47445 Moers
Germany

Phone: +49 941 29010-0
Fax: +49 941 29010-50
info@nal-vonminden.com
www.nal-vonminden.com

CEO:
Sandra von Minden
Roland Meißner
Thomas Zander

Bank: Sparkasse Regensburg (GER-
MANY)
Kto.: 8400 170 73 BLZ.:750 500 00
IBAN: DE9875 0500 0008 4001 7073
BIC/SWIFT: BYLADEM1RBG

Commercial register::
Kleve, HRB 5679
VAT-ID: DE189 016086
Tax-No: 244/133/00130

Rapid Tests

Laboratory Diagnostics

Laboratory Service

Consulting & Service

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