

EC-Declaration of Conformity

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

product number: 243203N-20
product name: NADAL® COVID-19 Ag+Influenza A/B 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 2022-10-06

Moers,07.10.2020



Sandra von Minden
CEO
nal von minden GmbH