

CE

April 2020

20

EC Declaration of Conformity

We,

Labcon, North America

3700 Lakeville Highway, Petaluma, California, USA 94954

+1 707 766 2100, www.labcon.com

Declare with sole responsibility, that our product (s)

EDMA Class 24 09 Micro Biology Disposables as:

Pipet Tips (Pipette tips)

Centrifuge Tubes and (Micro) centrifuge tubes

Culture Tubes

Disposable Laboratory Plastics

PCR Disposables

EDMA Class 21 01 1001 CH Hardware and Assessoris as:

POC Blood Dispenser

Blood Slide Prep Tool

Meet the essential requirements of Council Directive 98/79/EC pertaining to in-vitro diagnostics. Pathway of conformity per Annex III.

The product(s) identified above meet requirements of the IVDD by meeting the following standards:

ISO 9001:2015 Quality Management Systems

ISO 14971:2007 Risk Assessment

EN980:2008 Symbols Used in Labeling Medical Devices

EN ISO 18113-1:2011 Labeling of In-Vitro Diagnostic Devices

EN ISO 18113-1:2011 ISO 15223-1:2007 Symbols to be used with In-Vitro Diagnostic Devices

Our authorized representative within the European community as explicitly defined in Article 1. § 2(g) of Directive 98/79 EEC is:

mdi Europa GmbH.

Langenhagener Straße 71,
D-30855 Langenhagen, Germany

+49 511 39089530

www.mdi-europa.com



CE

labcon

20