EC Certificate

mdc medical device certification GmbH
Notified Body 0483
herewith certifies that

ravo Diagnostika GmbH
Oltmannsstraße 5
79100 Freiburg
Germany

for the scope

in vitro diagnostic devices for determination of antibodies specific for toxoplasmosis
ToxoTool M-I ISAGA
ToxoTool A-I ISAGA
ToxoMultiTool ISAGA
ToxoTool DIFA

has introduced and applies a

Quality System

for the design, manufacture and final inspection.

The mdc audit has proven that this quality system meets all requirements according to

Annex IV – excluding Section 4 and 6 of the Council Directive 98/79/EC

of the European Parliament and of the Council of
27 October 1998 on in vitro diagnostic medical devices.

The surveillance will be held as specified in Annex IV, Section 5.

Valid from	2020-06-13
Valid until	2024-05-26
Registration no.	D1052800014
Report no.	P19-01794-165362
Stuttgart	2020-06-12

Head of Certification Body

medical device certification mdc

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-247.10.05

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