CERTIFICATE OF REGISTRATION

MedNet EC-REP GmbH
Borkstraße 10
48163 Münster
Germany

in its function of the European Authorized Representative, in accordance with the In Vitro Diagnostic Directive 98/79/EC, hereby confirms the registration of the following in vitro diagnostic medical device(s) into the German DIMDI data base

Quiz SARS-CoV-2-NAT System (Kit)
DIMDI Registration Number DE/CA22/1311-710.1-IVD
Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System: POCHE
DIMDI Registration Number DE/CA22/1311-711.1-IVD
Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System: Replacement Kit: Positive Control
DIMDI Registration Number DE/CA22/1311-715.1-IVD
Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System: Replacement Kit: LAMP Beads
DIMDI Registration Number DE/CA22/1311-714.1-IVD
Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System: Replacement Kit: PCR Biochip
DIMDI Registration Number DE/CA22/1311-713.1-IVD
Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System (App)
DIMDI Registration Number DE/CA22/1311-712-IVD
Class: Other device (all devices except Annex II and self-testing devices)

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Local Reg. Number HRB 17791 | Managing Directors: Matthias Heinz, Ole Stein
Bank Account: Vereinigte Volksbank Münster eG | IBAN: DE97 4036 1906 7233 1271 00 | BIC: GENODEM1IBB

Quiz SARS-CoV-2-NAT System: OS Buffer
DIMDI Registration Number DE/CA22/1311-1241-IVD
Class: Other device (all devices except Annex II and self-testing devices)

on behalf of

Cell ID Pte Ltd.
3 Gambas Crescent, Nordcom One, #09-08, Singapore 757088

according to the directive 98/79/EC of the European Parliament and of the Council of the European Union relating to *in vitro* diagnostic medical devices.

Münster, 14.01.2022

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Birthe Harms
on behalf of MedNet EC-REP GmbH

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