

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-IVD

Class: Other device (all devices except Annex II and self-testing devices)

Quiz SARS-CoV-2-NAT System: POCHE, White and Black

DIMDI Registration Number DE/CA22/1311-711-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-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-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, 17.02.2021



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