CERTIFICATE OF REGISTRATION

MedNet 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

COVID-19 Antigen Rapid Test Device (Integrated Throat Swab),
Other device (all devices except Annex II and self-testing devices), Registration No.:
DE/CA22/419-1111-IVD

on behalf of

BMT Biomarketing, Ltd
14 Hailan St., Or-Akiva, 3060000
Israel

in Germany

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, 14.01.2021

MEDNET
Network to Market
MedNet GmbH · Borkstraße 10 · 48163 Münster
Tel. +49 (0) 251 32266-0 · Fax: +49 (0) 251 3[illegible]
www.mednet-europe.com · info@mednet-[illegible]

i.A.
on behalf of MedNet GmbH