CERTIFICATE OF REGISTRATION

MedNet EC-REP GmbH
Borkstraße 10
48163 Münster
Germany

in its function of the European Authorized Representative, in accordance with the Regulation (EU) 2017/746 on *in vitro* diagnostic medical devices of the European Parliament and of the Council dated 5 April 2017 in its latest version and including its amendments, hereby confirms the registration of the following *in vitro* diagnostic medical devices into the German DMIDS data base

QuantGene 9600 Fluorescent Quantitative Detection System, class A

on behalf of

Hangzhou Bioer Technology Co., Ltd.
1192 BinAn Rd., Binjiang District,
Hangzhou, 310053, China

according to the Regulation (EU) 2017/746 of the European Parliament and of the Council of the European Union relating to *in vitro* diagnostic medical devices.

Münster, 12.08.2022

Dr. Eva Steffens
on behalf of MedNet EC-REP GmbH

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