

## 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

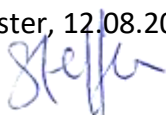
### **QuantGene 9600 Fluorescent Quantitative Detection System, class A**

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



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