Caretechion GmbH • Niederrheinstr. 71, 40474 Düsseldorf

开泰 启康 Caretechion

Guangzhou Decheng Biotechnology Co., Ltd.

Room 218, Building 2, No.68, Nanxiang Road, Science City, Huangpu District

510000 Guangzhou, China

11.12.2020

Certificate of Notification

This is to certify that, in accordance with the In Vitro Diagnostic Directive 98/79/EC, Caretechion GmbH agrees to as the EU Authorized Representative for:

Guangzhou Decheng Biotechnology Co., Ltd.

Room 218, Building 2, No.68, Nanxiang Road, Science City, Huangpu District

510000 Guangzhou, China

and confirms the submission of the notification of the *in vitro* diagnostic medical devices (as shown in Table 1) into the German DIMDI database according to The Act on Medical Devices (Gesetz über Medizinprodukte - MPG) of Germany.

The Manufacturer has provided Caretechion GmbH with the Declaration of Conformity confirming that the *in vitro* diagnostic medical devices fulfill the applicable requirement of In Vitro Diagnostic Directive 98/79/EC. In compliance with The Act on Medical Devices (Gesetz über Medizinprodukte - MPG) of Germany, a safety officer has been appointed..

Note: This certificate will automatically be invalid if the notification is canceled by the competent authority or exceeds the service scope or time of the EU Authorized Representative Agreement.

Product Name	DIMDI Form Number
2019-nCoV Ag Saliva Rapid Test Card (Immunochromatography)	00159870
2019-nCoV Ag Salvia Rapid Test Cup (Immunochromatography)	00159876

Table 1

With thanks and best wishes,

Caretechion GmbH
Niederrheinstr. 71, 40474 Düsseldorf
Tel.:0211 300 366 18 Fax:0211 300 366 19
E-mail: info@caretechion.de
www.caretechion.de
Amtsgericht Düsseldorf HRB 82833
Ust.-ID :DE319481632