Certificate
of EU product notification

Herewith we confirm that

MT Promedt Consulting GmbH
Altenhofstraße 80
66386 St. Ingbert
Germany

has taken over the function of an European Authorized Representative according to the requirements of Article 10 of the IVDD 98/79/EC for

RapiGen Inc.
2F, 25 Heungan-daero,
Gunpo-si, Gyeonggi-do 15809
Republic of Korea

MT Promedt Consulting GmbH has made the product notification at the relevant competent authority according to Article 10(3).
The in vitro diagnostic medical devices of the manufacturer, covered by the notification, are listed in Annex I of this certificate.

2 April 2020

Dr. Michael Rinck
- Managing Director -

Enclosure
Annex I

RapiGen Inc.

Annex I
to "Certificate of EU Product Notification"
(List of CE marked Products)

Internal Number	**Registration Number** (at the German CA/ DIMDI)	**Product Category (EDMS)**	**EDMS Code Description**	**Classification Annex**
RAP-01	DE/CA70/40838-154804	15 70 90 90 00	Other Other Virology Rapid Tests	III
RAP-02	DE/CA70/40838-135724	15 02 01 40 00	Hepatitis A Virus - NA Reagents	III
RAP-03	DE/CA70/40838-137534	15 70 01 02 00	H. Pylori - Rapid Test	III
RAP-04	DE/CA70/40838-139092	15 70 90 02 00	RSV - Rapid Test	III
RAP-05	DE/CA70/40838-139098	15 04 80 01 00	Adenovirus	III
RAP-06	DE/CA70/40838-139099	15 04 80 04 00	Influenza & Para Influenza	III
RAP-07	DE/CA70/40838-143028	15 70 01 05 00	Syphilis Rapid Tests	III
RAP-08	DE/CA70/40838-143177	15 04 80 06 00	Rotavirus	III
RAP-09	DE/CA70/40838-150810	12 70 13 03 00	Troponin I/T – Rapid Test	III

2 April 2020

Dr. Michael Rinck
- Managing Director -