DECLARATION OF NOTIFICATION

Date: March 17, 2020

The undersigned, Teresa Batet i Solà, Senior Consultant, of Qarad EC-REP BV, hereby declares that:

Shenzhen Bioeasy Biotechnology Co., Ltd.

No. 2-1, Liuxian 1st Road, Xin'an Sub-District, Baoan District,
Shenzhen, Guangdong Province, China. 518101

has signed the EC Declaration of Conformity in agreement with the Annex III of the European Directive 98/79/EC on In Vitro Diagnostic Medical Devices and has submitted the required technical documentation, for the following IVD products (for professional use only):

Name Device	Catalogue number Device
BIOEASY™ 2019-Novel Coronavirus (2019-nCoV) Ag GICA Rapid Test	YRLG22201025, YRLG22201050, YRLG22201100
BIOEASY™ Diagnostic Kit for 2019-Novel Coronavirus (2019-nCoV) Ag Test Kit (Fluorescence Immunochromatographic Assay)	YRLF04401025, YRLF04401050, YRLF04401100
BIOEASY™2019-Novel Coronavirus (2019-nCoV) IgG/IgM GICA Rapid Test (Colloidal Gold):	YRLG22301025, YRLG22301050, YRLG22301100
BIOEASY™ 2019-Novel Coronavirus (2019-nCoV) Ab GICA Rapid Test	YRLG22501025, YRLG22501050, YRLG22501100

The notification to the Belgian Competent Authorities has been carried out on March 12th, 2020 by Qarad EC-REP BV, the appointed Authorized Representative of Shenzhen Bioeasy Biotechnology Co., Ltd.

Information on the notification to the competent Authorities of other European countries is available upon request.

Teresa Batet i Solà
Senior Consultant

Qarad EC-REP BV
Authorized Representative