Declaration of Conformity

according to Directive 98/79/EC, on in vitro diagnostic medical devices

Ref. No.:20210125-E01

Maker (Name, Address)	**Getein Biotech, Inc.** No. 9 Bofu Road, Luhe District, Nanjing, 211505, China
Authorized Representative (Name, Address)	**Lotus NL B.V.** Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands.
Medical device	**Product Name** Getein1100 Immunofluorescence Quantitative Analyzer Getein1600 Immunofluorescence Quantitative Analyzer SARS-CoV-2 Total Antibody/Neutralizing Antibody Fast Test Kit (Immunofluorescence Assay) SARS-CoV-2 Neutralizing Antibody Fast Test Kit (Immunofluorescence Assay) One Step Test for SARS-CoV-2 Total Antibody/Neutralizing antibody (Colloidal Gold)
Classification	Others

Applicable coordination standards			
	EN 13612:2002	EN ISO 14971:2012	EN ISO15223-1:2016
	EN ISO 18113-1:2011	EN ISO 18113-2:2011	EN ISO 18113-3:2011
	EN ISO 23640:2015	EN ISO 13485:2016	ISO 780:2015
	EN 61326-2-6:2006	IEC 61326-1:2013	
	EN 61010-2-101:2002	IEC 61010-1:2010	

Signatory representative declares herein the above mentioned device meets the basic requirements of the European Parliament and the Council's in vitro diagnostic medical devices directive: 98/79/EC Annex III. This declaration of conformity is based on European Parliament and the Council's 98/79/EC directive Annex III. The compiled technical file and quality system document according to 98/79/EC directive Annex III are testified and the quality system certificate has issued by BSI Group The Netherlands B. V..

General Manager Enben Su

Nanjing, 25th Jan. 2021

(place and date of issue)

Enben Su

(name and signature or equivalent marking of authorized person)

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