## **EC Declaration of Conformity**

Manufacturer:

Name: JOYSBIO (Tianjin) Biotechnology Co., Ltd.

**Address:** Tianjin International Joint Academy of Biotechnology & Medicine 9th floor, No.220, Dongting Road, TEDA 300457 Tianjin

China.

**Tel:** +86-022-65378415 **Email:** molly@joysbio.com

We, the manufacturer, here with declare that the product(s)

## **Whose Authorized Representative:**

Name: Lotus NL B.V.

**Address:** Koningin Julianaplein 10,1e Verd, 2595AA, The Hague, Netherlands.

E-mail: peter@lotusnl.com

Product Name	SARS-CoV-2 Antigen Rapid Test Kit	Specification	20Tests/box (1Test/bag
	(Colloidal Gold)		×20 Bags)
	For in vitro qualitative detect of SARS-CoV-2 nucleocapsid antigen in saliva specimens directly from		
Intended Use	individuals who are suspected of COVID-19 by their healthcare provider within the first 5 days of the		
	onset of the symptoms. This test is only provided for use by clinical laboratories or to healthcare workers		
	for point-of-care testing, and not for at home testing.		
Classification	Others		
Classification	for point-of-care testing, and not for at home testing.	~)///((////////////////////////////////	

Conformity Assessment Route: IVDD98/79/EC Annex III.

## **Applicable Standards:**

ISO 13485:2016 ISO 14971:2019 EN ISO 18113-1:2011 EN ISO 18113-2:2011 EN ISO 18113-3:2011 EN 13641:2002 ISO 15223-1:2016 EN 13612:2002 ISO 23640:2015 EN 62366-1:2015



We, the manufacturer, here declare with sole responsibility that our product/s mentioned above meet/s the provisions of the Directive 98/79/EC of the European Parliament and of the Council on In-Vitro Diagnostic Medical Devices.

We agree to develop, implement and maintain a documented post-production monitoring process.

Nout ben 10/11/2020	
10/11/2020	
10/11/2020	
Tianjin, China Biotecho	
正元成邦(天津)	
生物科技有限公司	