

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

Whose Authorized Representative:

Name: Lotus NL B.V.
Address: Koningin Julianaplein 10,1e Verd, 2595AA, The Hague, Netherlands.
E-mail: peter@lotusnl.com

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

| | | | |
|-----------------------|--|----------------------|---|
| Product Name | SARS-CoV-2 Antigen Rapid Test Kit (Colloidal Gold) | Specification | 20Tests/box (1Test/bag ×20 Bags) , 40 Tests /box (1Test / bag ×40 Bags) |
| Intended Use | For in vitro qualitative detect of SARS-CoV-2 nucleocapsid antigen in nasal(NS) swab specimens directly from 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 | | |

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

Applicable Standards:

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



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.

| | |
|--------------------------------|-----------------|
| Name of General Manager | 王森 |
| Signature | |
| Date | 2020.08.28 |
| Place | Tianjin, China. |
| Seal (Manufacturer) | |