

## Certificate of Compliance

This is to certify that the following designated products:

ELISA & RAPID TEST KITS FOR HUMAN, VETERINARY & PLANT RESEARCH & DIAGNOSTICS

## **Manufactured By**

## ARSH BIOTECH PRIVATE LIMITED

308, Third Floor, Aggarwal City Mall, Opposite M2K, Pitampura,

**Delhi - 110034, India** 

Conforms with the Applicable requirements of

Vitro Diagnostic Medical Devices (98/79/EC)

Certificate No.: 20HR125C

## This certificate is issued under the following conditions:

- 1. It applies only to the above referenced set of products mentioned above. The manufacturer is obligated to assure that all products of the respective model confirm to the type assessed by this certificate.
- 2. The Certificate validity is conditioned by the positive result of the surveillance audits.
- 3. The Certificate remains valid until the manufacturing conditions, the quality systems or relevant legislation are changed but until the 13/01/2024.
- 4. After Fulfilling the EU legislation requirements, the manufacturer shall affix to each product of the above referenced models, CE Marking according to the following example.

Issued Date: 14/01/2021 1st Surveillance: 13/01/2022

Expiry Date: 13/01/2024 2nd Surveillance: 13/01/2023







INTERNATIONAL QUALITY REGISTRAR CERTIFICATION

Second Floor, 1 Berkeley Street, W1J 8DJ London (UK)

This Certificate is intellectual Property of IQRC and can be maintained through surveillance and renewal adults

Certificate should be returned to IQRC in case of non compliance of certification procedure

Authenticity of this certificate can be verified at www.ukqc.org.uk

The Registration is not a Product Quality Certificate.