

Certificate of Registration

This is to Certify that
Quality Management System of

YUVRAJ BIOBIZ INCUBATOR INDIA PRIVATE LIMITED

**W - 2, WATER WORKS ROAD, THIRU VI KA INDUSTRIAL ESTATE,
GUINDY, CHENNAI – 600032, TAMIL NADU, INDIA**

has been assessed and found to conform to the requirements of

ISO 9001:2015

for the following scope :

**DESIGN, MANUFACTURE AND SUPPLY OF IN VITRO DIAGNOSTIC
MEDICAL DEVICES SUCH AS RAPID TEST KITS, ELISA TEST KITS,
BIOCHEMISTRY REAGENTS, SEROLOGY REAGENTS, HAEMATOLOGY
REAGENTS AND POINT OF CARE DEVICES**

Certificate No	: 21IQHF36	Issuance Date	: 19/07/2021
Initial Registration Date	: 19/07/2021		
Date of Expiry	: 18/07/2024		
1st Surve. Due	: 19/06/2022	2nd Surve. Due	: 19/06/2023


Director



AQC MIDDLE EAST LLC

Head Office: Office No. 02, Ground Floor, Sharjah Media City, Sharjah, UAE. e-mail : info@aqcworld.com,

Key Location: A-60, Sector - 2, Noida, Uttar Pradesh, 201301, India.

*Validity of the Certificate is subject to successful completion of surveillance audit on or before of due date. (in case surveillance audit is not allowed to be conducted, this certificate shall be suspended/withdrawn).

Certificate Verification: Please Re-check the validity of certificate at <http://www.aqcworld.com/activeclients.aspx> or www.aqcworld.com at Active Clients.

Certificate is the property of AQC Middle East LLC and shall be returned immediately when demanded



ACCREDITED
Management Systems
Certification Body
MSCB-119



ISO 9001:2015





Ref. No. 21IQHF36/S1

Dated: 01/06/2022

To,

YUVRAJ BIOBIZ INCUBATOR INDIA PRIVATE LIMITED
W - 2, WATER WORKS ROAD, THIRU VI KA INDUSTRIAL ESTATE, GUINDY,
CHENNAI – 600032, TAMIL NADU, INDIA.

Subject: Continuation of ISO 9001: 2015 Certificate

Dear Sir,

Please accept our congratulations on successful completion of 1st Surveillance Audit for Quality Management System Certification at your works/Office for continued compliance towards the requirements of ISO 9001:2015 standard.

Scope of Registration;

**DESIGN, MANUFACTURE AND SUPPLY OF IN VITRO DIAGNOSTIC MEDICAL
DEVICES SUCH AS RAPID TEST KITS, ELISA TEST KITS, BIOCHEMISTRY
REAGENTS, SEROLOGY REAGENTS,
HAEMATOLOGY REAGENTS AND POINT
OF CARE DEVICES**

Certificate No. – 21IQHF36/R1

The Certificate will remain valid till: **18/07/2024**

The Second Surveillance Audit need to be conducted by or before: **19/06/2023**

Regards,

Director