

CERTIFICATE OF REGISTRATION

INTERCERT hereby certifies that the Quality Management System of



Central Drug House (P) Ltd

Corp Office: 7/28 Vardaan House, Darya Ganj, New Delhi - 110002, India **Manufacturing Unit:** D-2/CH/9 Dahej-2 GIDC,Bharuch - 392130 Gujarat, India

Has been successfully assessed as per the requirements of

ISO 9001:2015

For the scope of Manufacturing, Marketing, Distribution, Technical Support for Laboratory Reagents, Fine Chemicals, Research Chemicals, Dehydrated Culture Media, Culture Media Bases, Filter Papers, Membrane Filters and Disinfectant

A	May 09, 2019	
:	May 09, 2022	
9	May 08, 2023	
:	May 08, 2025	
	ļ	 May 09, 2019 May 09, 2022 May 08, 2023 May 08, 2025

Registration Number: IC-QM-1905033

Issued on behalf of InterCert Head - Certifications



The validity of this certificate can be verified at **www.intercert.com** or through email at **info@intercert.com**. This certificate is the property of **INTERCERT INC**, 2001 Timberloch Place - Suite 500, The Woodlands, Texas 77380, United States and must be returned on request.



CERTIFICATE OF REGISTRATION

INTERCERT hereby certifies that the Environmental Management System of



Central Drug House (P) Ltd

Corp Office: 7/28 Vardaan House, Darya Ganj, New Delhi - 110002, India **Manufacturing Unit:** D-2/CH/9 Dahej-2 GIDC,Bharuch - 392130 Gujarat, India

Has been successfully assessed as per the requirements of

ISO 14001:2015

For the scope of

Manufacturing, Marketing, Distribution, Technical Support for Laboratory Reagents, Fine Chemicals, Research Chemicals, Dehydrated Culture Media, Culture Media Bases, Filter Papers, Membrane Filters and Disinfectant

Initial Certification Date	: May 09, 2019
Certificate Issue Date	: May 09, 2022
Surveillance Validity Date	: May 08, 2023
Recertification Date	: May 08, 2025

Registration Number: IC-EM-1905034



Issued on behalf of InterCert Head - Certifications



The validity of this certificate can be verified at **www.intercert.com** or through email at **info@intercert.com**. This certificate is the property of **INTERCERT INC**, 2001 Timberloch Place - Suite 500, The Woodlands, Texas 77380, United States and must be returned on request.



Certificate of Registration

This is to certify that The Quality Management System of Medical Devices



Corp Office: 7/28 Vardaan House, Darya Ganj, New Delhi - 110002, India Manufacturing Unit: D-2/CH/9 Dahej-2 GIDC, Bharuch - 392130 Gujarat, India

has been assessed and found to be in compliance with the requirements of the standard

ISO 13485:2016

for the following scope :

Manufacturing, Marketing, Distribution, Technical Support for Laboratory Reagents, Fine Chemicals, Research Chemicals, Dehydrated Culture Media, Culture Media Bases, Filter Papers, Membrane Filters and Disinfectant.

CERTIFICATE NO. : 19ZKCM02144M

INITIAL ISSUE DATE : 27/06/2019 REISSUE DATE **EXPIRY DATE**

27/06/2022 : 26/06/2025

1 ST SURVEILLANCE 26/06/2023 26/06/2024 2ND SURVEILLANCE :



Authorised Signatory

INTERNATIONAL QUALITY CERTIFICATION SERVICES UK LTD 272, Bath Street, Glasgow, G2 4JR, U.K.

This Certificate is intellectual Property of IQCS and can be maintained through surveillance and renewal audits. Certificate should be returned to IQCS in case of non compliance of certification procedure. Authenticity of this certificate can be verified at www.ukacert.co.uk / www.iqcscert.co.uk The Registration is not a Product Quality Certificate.







Certificate of Compliance

This is to Certify that the following designated products :

Laboratory Reagents, Fine Chemicals, Research Chemicals, Dehydrated Culture Media, Culture Media Bases, Filter Papers, Membrane Filters and Disinfectant.

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 survellance audits.
- 3. The Certificate remains valid until the manufacturing conditions, the quality systems or relevant legislation are changed but until the 26/06/2025
- 4. After Fulfilling the EU legislation requirments, the manufacturer shall affix to each product of the above referenced models, CE Marking according to the following example

Manufactured By

Central Drug House (P) Ltd

Corp Office: 7/28 Vardaan House, Darya Ganj, New Delhi - 110002, India Manufacturing Unit: D-2/CH/9 Dahej-2 GIDC, Bharuch - 392130 Gujarat, India

Conforms with the Applicable requirements of

In Vitro Diagnostics Directive (IVD) 98/79/EC

CERTIFICATE NO. : 19ZKCM02145C

INITIAL ISSUE DATE : 27/06/2019 REISSUE DATE **EXPIRY DATE**

27/06/2022 : 26/06/2025

1 ST SURVEILLANCE 2ND SURVEILLANCE •

26/06/2023 26/06/2024

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