By Royal Charter

# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 9001:2015

This is to certify that:

TCS Biosciences Limited
Botolph Claydon
Buckingham
MK18 2LR
United Kingdom

Holds Certificate Number: FS 28907

and operates a Quality Management System which complies with the requirements of ISO 9001:2015 for the following scope:

**The procurement, manufacture and sale of a range of diagnostic products for clinical, pharmaceutical, food and environmental laboratory testing.**

Digitally signed by Monastîrschii Viorica
Date: 2022.01.10 14:46:04 EET
Reason: MoldSign Signature
Location: Moldova

For and on behalf of BSI:

Andrew Launn, EMEA Systems Certification Director

Original Registration Date: 1994-08-11
Latest Revision Date: 2019-01-17

Effective Date: 2019-01-27
Expiry Date: 2022-01-26



...making excellence a habit.™

By Royal Charter

# Certificate of Registration

## ENVIRONMENTAL MANAGEMENT SYSTEM - ISO 14001:2015

This is to certify that:

TCS Biosciences Ltd
Botolph Claydon
Buckingham
MK18 2LR
United Kingdom

Holds Certificate Number: EMS 590359

and operates an Environmental Management System which complies with the requirements of ISO 14001:2015 for the following scope:

**The procurement, manufacture and sale of a range of diagnostic products for clinical, pharmaceutical, food and environmental laboratory testing.**

For and on behalf of BSI:

Andrew Launn, EMEA Systems Certification Director

Original Registration Date: 2013-06-10
Latest Revision Date: 2019-01-18

Effective Date: 2019-01-27
Expiry Date: 2022-01-26



...making excellence a habit.™

UKAS MANAGEMENT SYSTEMS 003

## SELF DECLARATION OF CONFORMITY

We declare under our sole responsibility in accordance with MHRA Registration Number IVD 000100 that the following CE marked products:

| EDMA code(s) | EDMA description | TCS product code and description |
|---|---|---|
| 14.50.01.90 | Other Controls/Standards/Calibrators, Microbiology | Selectrol - All MM codes |

conform to the relevant provisions of the In-vitro Diagnostic Medical Devices Directive 98/79/EC and The Medical Devices Regulations 2002 (SI 2002 No.618) and The Medical Devices (Amendment) Regulations 2003 (SI 2003 No.1697) for in-vitro diagnostic medical devices.

This declaration is made on the basis of meeting the requirements of Annexes I and III of the In-Vitro Diagnostic Medical Devices Directive 98/79/EC and continued maintenance of an approved Quality Management System meeting the requirements of ISO 9001, as certified by BSi, certificate number FS 28907.

Signed by: Sue Brown  Date: 30.04.2016

Name: Sue Brown
Position: Regulatory Affairs Manager

Signed by: Lynda Preston  Date: 30.04.2016

Name: Lynda Preston
Position: Managing Director