

## **CERTIFICATE OF REGISTRATION**

This is to certify that the management system of:

## **Oxoid Company**

(FIN F006846) Main Site: 1926 Merivale Road, Unit 100,

Nepean, Ontario, K2G 1E8, Canada

Additional Site: 1886 Merivale Road,

Nepean, Ontario, K2G 1E8, Canada

has been registered by Intertek, an MDSAP recognized auditing organization, as conforming to the requirements of:

## ISO 13485:2016

Canada: Medical Devices Regulations – Part 1- SOR 98/282

## The management system is applicable to:

Design/development, manufacture and distribution of the following in-vitro diagnostic medical devices used in the isolation, identification and susceptibility testing of micro-organisms; - ready-prepared, prepoured microbiological non-sterile culture media.

Main site: Management, Quality, Design/development transfer, Manufacturing, Storage, and Purchasing

Additional site: Storage and Distribution.

Certificate Number: 0149296-02

Initial Certification Date: 2021-12-17

**Certification Effective Date:** 2024-12-16

**Certification Expiry Date:** 2027-12-16



intertek

Calin Moldovean President, Business Assurance

Intertek Testing Services NA, Inc. 4700 Broadmoor SE, Suite 200 Kentwood, MI, USA, 49512





In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone. The certificate remains the property of Intertek, to whom it must be returned upon request. Validity of this certificate may be verified at <a href="http://www.intertek.com/business-assurance/certificate-validation/">http://www.intertek.com/business-assurance/certificate-validation/</a>

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