By Royal Charter

# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Curiteva, Inc
25127 Will McComb Drive NW
Tanner
Alabama
35671
USA

Facility ID Number: F005720

Holds Certificate No: **MDSAP 740824**

The company listed on this certificate has been audited to and found to conform with ISO 13485:2016 including the following country specific requirements:

**USA:** 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

Design, development and manufacture of implantable orthopedic hardware including sterile and non-sterile cervical interbody fusion implants and non-active surgical instruments from titanium, stainless steel, PEEK and like materials.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2021-11-19 Effective Date: 2024-07-30 Expiry Date: 2027-07-29



MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM
BSI Group America Inc. is an MDSAP recognised auditing organization

...making excellence a habit.™

This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated online. Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: BSI Group America Inc., 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA
A Member of the BSI Group of Companies.