By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Anika Therapeutics, Inc.
32 Wiggins Ave.
Bedford
Massachusetts
01730
USA

Facility ID Number: F009203

Holds Certificate No: **MDSAP 845340**

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

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure
Brazil: RDC ANVISA n. 67/2009, RDC ANVISA n. 665/2022 - Good Manufacturing Practices, RDC ANVISA n. 551/2021
Canada: Medical Devices Regulations - Part 1 - SOR 98/282
USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

Please see scope page.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2026-07-06 Effective Date: 2026-07-06 Expiry Date: 2028-03-28

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.

America Headquarters: BSI Group America Inc. 1950 Opportunity Way, Suite 900, Reston, VA 20190.
A Member of the BSI Group of Companies.

Certificate No: **MDSAP 845340**

Registered Scope:

Design and manufacture of sterile hyaluronic acid for surgical, orthopaedic, ophthalmic application.

Design and development, manufacture and distribution of ancillary Instruments for the Area of orthopaedic Surgery.

Certificate transfer from GMED
Certificate number: N° 36681–7

Original Registration Date: 2026-07-06 Effective Date: 2026-07-06 Expiry Date: 2028-03-28

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