By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Micropore, Inc.
1000 Konica Drive
Elkton
Maryland
21921
USA

Facility ID Number: F008249

Holds Certificate No: **MDSAP 827258**

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
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

Design and manufacture of carbon dioxide absorbents and absorbent canisters for the area of anesthesiology.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2026-04-27 Effective Date: 2026-04-27 Expiry Date: 2029-04-26

MDSAP
Medical Device Single Audit Program
BSI Group America Inc. is an MDSAP recognised auditing organization

...making excellence a habit.™

Certificate No: **MDSAP 827258**

Location	Registered Activities
Micropore, Inc. 1000 Konica Drive Elkton Maryland 21921 USA Facility ID Number: F008249	Management activities, design, extraction, finishing sorbents and canisters and final testing.
Micropore, Inc. 350F Pencader Drive Newark Delaware 19702 USA Facility ID Number: F008249	Extrusion and extraction processes.

Original Registration Date: 2026-04-27 Effective Date: 2026-04-27 Expiry Date: 2029-04-26