

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Ribbond Inc
1904 3rd Ave, Suite 330
Seattle
Washington
98101
USA

Facility ID Number: F001453

Holds Certificate No:

MDSAP 699174

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

Japan: MHLW MO No 169 (2004), as amended by MHLW MO No 60 (2021), PMD Act

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

The design, manufacture and distribution of dental reinforcement material, including the Starter Kit and the Refill, and of dental resins.

For and on behalf of BSI:



Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2020-10-02

Effective Date: 2023-07-11

Expiry Date: 2026-07-10



BSI Group America Inc. is an MDSAP recognised auditing organization

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