



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that: DJO FRANCE SAS

Centre Européen de Frêt

3 rue de Bethar Mouguerre 64990 France

Facility ID Number: F000514

Holds Certificate No: MDSAP 695371

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

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

Please see scope page.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2019-07-18 Effective Date: 2024-08-30 Expiry Date: 2024-05-21

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MEDICAL DEVICE SINGLE AUDIT PROGRAM

BSI Group America Inc. is an MDSAP recognised auditing organization

...making excellence a habit."

Certificate No: MDSAP 695371

Registered Scope:

Design, manufacture, distribution and service of Radial Pressure Wave devices, Neuromuscular stimulators, Ultrasound units and their software.

Design, manufacture and distribution of Iontophoresis system electrodes and non-sterile accessories.

Distribution of orthopaedic units and diathermy units.



Original Registration Date: 2019-07-18 Effective Date: 2024-07-05 Expiry Date: 2027-05-21

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