



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that: Össur Iceland ehf.

Grjótháls 5 Reykjavik 110 Iceland

Facility ID Number: F001444

Holds Certificate No: MDSAP 813028

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

The design, manufacture and distribution of external prosthetic devices, orthotic devices and rehabilitation circulation packs. The service of external prosthetic devices and orthotic devices.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2025-08-05 Effective Date: 2025-12-16 Expiry Date: 2028-12-15

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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 813028

Location Registered Activities

Össur Iceland ehf. Grjótháls 5 Reykjavik 110 Iceland

Facility ID Number: F001444

Ossur Americas 680 Grove Rd. West Deptford New Jersey 08086 USA

Facility ID Number: F003629

The design, manufacture and distribution of external prosthetic devices, orthotic devices and rehabilitation circulation packs. The service of external prosthetic devices and orthotic devices.

The manufacture, distribution, and service of external prosthetic devices. The manufacture and distribution of orthotic devices. The distribution of rehabilitation circulation packs.



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