

# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

NeoCoil, LLC  
N27 W23910A Paul Road  
Pewaukee  
Wisconsin  
53072  
USA

Facility ID Number: F000329

Holds Certificate No:

**MDSAP 689260**

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

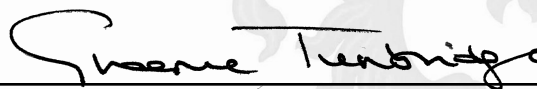
**Canada:** Medical Devices Regulations - Part 1 - SOR 98/282

**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, development, manufacture, servicing, installation and distribution of magnetic resonance radio frequency coils and accessories for diagnostic imaging in the medical industry.

For and on behalf of BSI:



Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2019-05-24

Effective Date: 2025-03-24

Expiry Date: 2028-03-23



BSI Group America Inc. is an MDSAP recognised auditing organization

Page: 1 of 1

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