



Norman Noble, Inc. Announces ISO 13485:2016 Certification

HIGHLAND HEIGHTS, OHIO – December 12, 2018 – Norman Noble, Inc., the world’s leading contract manufacturer of next-generation medical implants, has achieved ISO 13485:2016 registration for its Quality Management System (QMS) following a full recertification audit by the British Standards Institute (BSI), a European Union Notified Body. ISO 13485:2016 is an internationally recognized quality standard specific to the medical device industry. ISO 13485:2016 registration demonstrates Norman Noble’s continued commitment to the highest level of medical device product quality and regulatory compliance. ISO 13485 registered since 2004, this update supports our customers’ efforts to maintain compliance with the latest domestic (FDA) and international (MDSAP and European MDR) regulatory requirements.

About Norman Noble

Established more than 70 years ago, Norman Noble, Inc. remains a family-owned and -operated company offering the most advanced processes for ultra-precision micromachining of medical implants and aerospace parts. The company is known for its exceptional ability to produce nitinol based implants, and achieve sub-miniature precision beyond the reach of most manufacturers. Norman Noble is a supplier to most of the largest OEM’s and well-known names in the medical device industry.

Norman Noble manufactures medical devices and implants to customer specifications in compliance with FDA regulations and ISO 13485. State-of-the-art processes include athermal laser machining, laser welding, Swiss turning and milling, conventional and wire EDM, high-speed 7-axis contour milling, electropolishing, nitinol shape setting and clean room assembly and packaging. Rapid development prototyping services are available in separate and fully dedicated process development centers. FDA Registration #1531050. Virtual tour and more information: www.nnoble.com.

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