4 January 2024

NEWS

Ferrosan Medical Devices A/S Achieves Key Milestone with First MDR Certifications

We are pleased to announce that Ferrosan Medical Devices A/S has been awarded the first two MDR certifications, marking a significant milestone and demonstrating our commitment to delivering high-quality medical devices that meet the latest European Union's Medical Device Regulation (MDR), which places stringent requirements on the production and distribution of medical devices.

SURGIFLO™ Haemostatic Matrix (MS0010) and SURGIFLO™ Haemostatic Matrix Kit with Thrombin (MS0012) are now certified according to the new European Medical Device Regulation (EU) 2017/475 on Medical Devices (MDR). The MDR certificate gives Ferrosan Medical Devices A/S the right to continue to market Flowables within the European Union and facilitate registrations outside the European Union as it has been adapted as a guiding certificate by many countries.

About Ferrosan Medical Devices

Ferrosan Medical Devices develops and manufactures medical devices used in surgical procedures by health care professionals all over the world. Every two seconds a device from Ferrosan Medical Devices is used. We have a mission to provide innovative, effective, and safe medical devices that enable surgeons, nurses, and clinicians to perform surgical procedures as seamlessly as possible without complications. Making seconds count in surgical care.
The products are sold and marketed under a global partnership agreement with Ethicon, Inc. (part of Johnson & Johnson). In addition, FeMD develops and produces handheld biopsy devices, used in the diagnosis of breast cancer. These are developed and distributed in collaboration with a global partner. FeMD has around 350 employees and production facilities in Søborg in Denmark and Szczecin in Poland.