Beaufort®

Accelerating Medical Innovation

Are You Ready for MDR?

The time for action is now!
The Beaufort MDR Team Can Help

With May, 2020 just around the corner, Beaufort is assisting sponsors with comprehensive services to optmize the transition from MDD to MDR.

Our MDR subject matter experts understand the complex details, regulatory expectations and new mandatory requirements. We will design and implement a comprehensive MDR Action Plan for your portfolio of medical devices for the EU market, which can include:

- MDR Services
  - Clinical data and documentation
  - Quality system
  - Product classification
  - Technical documentation
  - Unique Drive Identification (UDI)
  - Post-market surveillance
  - ISO 13485:2016
- Remediation and Implementation
  - Technical file and report updates
  - Clinical literature searches and data consolidation
  - Clinical trial planning and management
- Notified Body Support

Our MDR team will provide the people, processes, and support needed to help you address the regulations.

About Beaufort CRO

Beaufort is an ISO 9001:2015 certified, global contract research organization that partners with medical device manufacturers, in vitro diagnostic (IVD), companion diagnostic, and clinical laboratories to provide a full range of clinical, regulatory, and quality solutions, including MDR expertise to meet new EU requirements.

For more than 15 years, companies worldwide have trusted Beaufort for its client-focused approach to research. Our deep industry expertise and tested and proven solutions have a track record of success in bringing our clients' products to market.

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