Regulatory Compliance Solutions
EU MDR

Raland Compliance Partners is your trusted partner delivering sustainable Quality, Regulatory, and Clinical compliance solutions.

Raland Compliance Partners (RCP) provides full life-cycle compliance consultation for **Medical Device**, Pharmaceutical, and Biotech companies worldwide.

With the 3-year implementation of the new Medical Devices Regulation (EU MDR) well underway, many manufacturers are finding themselves behind the curve in completing even the initial steps necessary to meet the May 2020 deadline and may be at risk of losing their CE Mark and access to the EU market.

The RCP Regulatory Compliance team is ready to lead or augment your internal team and provide the necessary guidance and manpower to catch up to the MDR timeline. Our team recommends the following programmatic methodology to accomplish the steps necessary to comply with EU MDR:

- Educate – promote understanding and the implications of the new regulation
- Organize – develop a comprehensive list of CE Marked products
- Assess – review and prioritize the product list
- Plan – develop plan for all phases of transition
- Execute – mobilize team to execute the plan
- Sustain – incorporate training and update processes and procedures

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