



**GLOBAL COMPLIANCE
MADE SIMPLE**



ASSURING PRODUCT COMPLIANCE AND MAINTAINING TRUST

At Cranage Veritas, we believe product compliance is far more than a checklist — it's a commitment to ensuring that innovative products are safe, positively impact society, and meet complex regulatory requirements. Safety is central to our work, and we support designers and manufacturers with consultancy, guidance, and advice across product safety, regulatory affairs, compliance, and market access to help ensure a smooth path from concept to a market-ready product.

Our team brings many years of experience across industries such as medical devices, IT, and electric vehicles, enabling us to deliver expert, comprehensive support throughout the design process. We assist manufacturers during early development, through detailed compliance planning, and into post-production assessments and market approval submissions, helping them achieve the standards required for successful market entry.





OUR SERVICES

Cranage Veritas provides a focused range of services to help manufacturers achieve compliance and gain market access. We support companies through regulatory requirements, product approvals, and representation in both the EU and UK.

Our services include:

- **Regulatory Affairs & Market Access** – Support with IEC 60601-1, EU MDR, and FDA 510(k) processes, including documentation and regulatory guidance.
- **EU/UK Representation** – Acting as an Authorised Representative and providing PRRC services for medical-device manufacturers.
- **CE Marking Support** – Identifying applicable directives, preparing technical files, and managing conformity assessments.
- **Local Technical Representative (LTR)** – Supporting CB Scheme certification through QMS reviews and witnessed factory testing.
- **UK Responsible Person** – Handling MHRA registrations, technical documentation, and post-market compliance for UK-market devices.

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