



MARKET ACCESS HAS NEVER BEEN EASIER

US AGENT
EU REPRESENTATIVE
UK RESPONSIBLE PERSON
SWISS REPRESENTATIVE
GDPR REPRESENTATIVE
DATA PROTECTION OFFICER



MEDITRIAL HAS HELPED MEDICAL DEVICE AND IVD COMPANIES WITH REGULATORY COMPLIANCE AND MARKET ENTRY SINCE 2008.

WITH OFFICES IN EUROPE AND THE US, MEDITRIAL OFFERS A WIDE RANGE OF MARKET ACCESS SERVICES, INCLUDING REGULATORY COMPLIANCE, CONSULTING, DEVICE REGISTRATION, AND IN-COUNTRY REPRESENTATION.

DEVICE REGISTRATION

We utilize years of hands-on experience to ensure our clients get their products to market as efficiently as possible.

IN-COUNTRY REPRESENTATION

Meditrial serves as a strategic partner, helping companies comply with legal requirements and establish constructive relationships with authorities, payors, hospital networks and academic associations.

POST-MARKET SURVEILLANCE

We provide post-market compliance solutions, incident reporting, and vigilance support driven by a life-cycle approach to device safety and effectiveness.

