IQVIA MedTech Real World Solutions

Device utilization electronic survey: Using RWE to address medical device regulation requirements in post-market surveillance

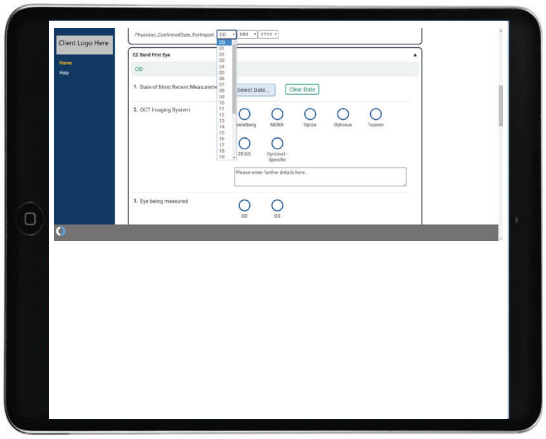
A simple method to capture post-market use of a medical device in the real-world setting, satisfying the need of EU MDR Article 83* to gather, record and analyse the ongoing performance of your product.

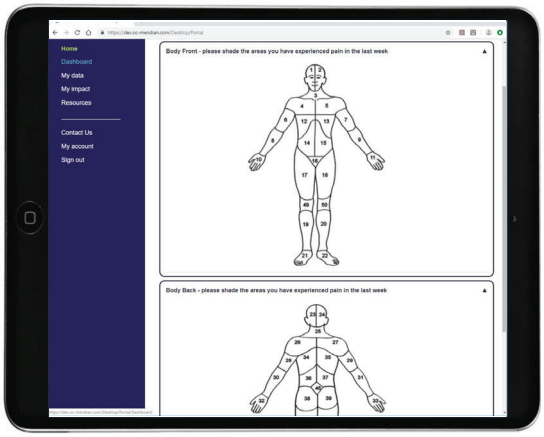
SCOPE

- Assess the real-world usage of medical devices
- Capture real world data for post-market surveillance
- Real-time access to data
- Provide on-demand reports
- Improve device usability, performance and safety

KEY FEATURES

- Maximize data generation
- Rapid set-up timelines
- Cost effective solution
- Ease of use for Physicians / Surgeons and Sponsor
- PRO integration option
- Physician/ Stakeholder consent capture option
- Supports multiple languages





*European Union Medical Device Regulation (Council Regulation 2017/745): Chapter VII, Section 1, Article 83

If you would like more information about this solution, please contact us via **iqviamedtech.com/rwe**