

Medical Device Registries - Bridging the Gap Between Device Performance in Clinical Trials And Their Use in Routine Practice

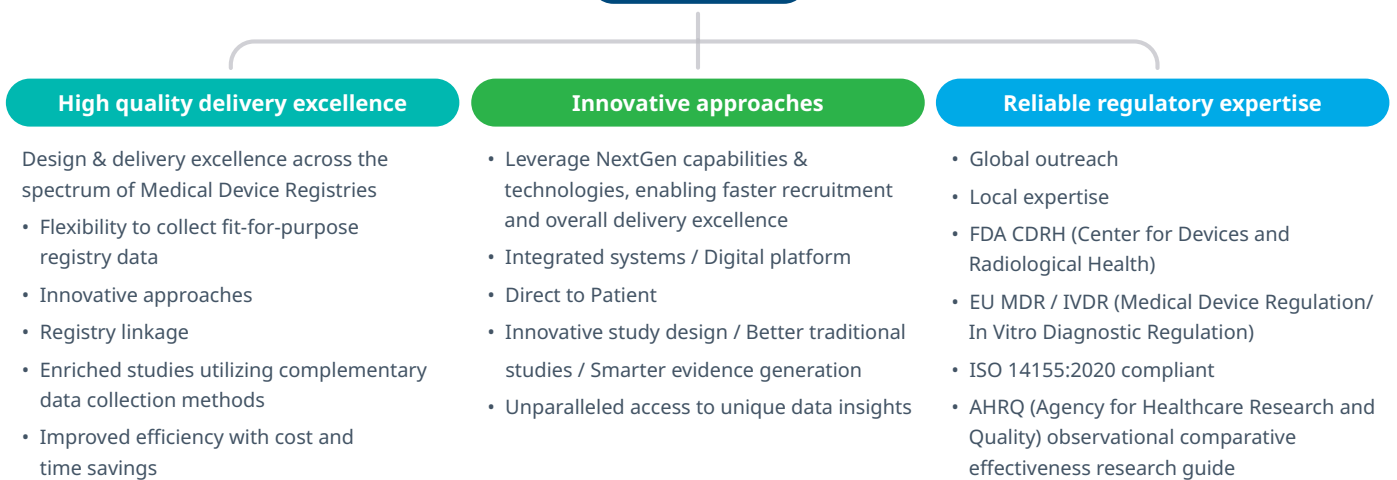
With the continuous evolution of the medical device regulatory landscape, especially with requirements to proactively demonstrate product safety, effectiveness and performance in a real-world setting, the need to build smart and cost-effective registries became a must-have.

Device registries allow assessment of **Medical device registries** performance in a real-world setting, to meet multi-stakeholder needs.

Medical device registries to meet multi-stakeholders needs



Why IQVIA?



CONTACT US

iqviamedtech.com