CLINICAL AFFAIRS

Bannick LLC
a primary device group

BannickPrimary.com

Bannick provides clinical, regulatory, quality, auditing and medical writing consulting services to the medical device industry to help you get your product through regulatory approvals and to market.

Clinical Affairs Consulting Services We Offer:

- Clinical study strategy and design
- Clinical validation
- Post-market clinical follow-up
- Protocol and CRF development
- Clinical study management
- Investigator meetings & site training
- Clinical monitoring
- Clinical audits

Optimize your path to regulatory approval by leveraging our unparalleled clinical affairs and development expertise. At Bannick, we understand the importance of timely, high-quality, and cost-effective clinical development. Our services ensure scientific integrity and strategic execution of deliverables.

Your clinical study is a critical element in your regulatory pathway. Our clinical experts can guide your company to compliance with FDA and the new EU Medical Device Regulations (MDR) requirements.