Protocol Next

AI-Driven Protocols for Smarter Trials

30 years of clinical research solution expertise

Description

Our **cloud-based** SaaS platform uses **AI and ML** to **analyze** data, **identify** patterns, and **predict** outcomes, **streamlining** protocol design while keeping your proprietary data **secure**. **Accelerate** development, **minimize** risks, and ensure built-in **compliance** and **quality**—all in one solution.

Your Challenges:

- Complex & Extensive data input
- Lengthy time for development
- Study design requires manual review of documents, trials, standards, patients' requirements, etc…
- Multiple templates and guidance's
- Tedious process for Unstructured data review

Our Solution:

- SaaS Based product with tiered approach
- Streamlined trial design using intelligent templates and deep historical analytics
- AI/ML synthesized trial content based on data input and study intent
- Automated data recommendations with alerts and notification for review and acceptance
- Model-driven treatment arms, randomization, drug dose calculations
- Verification of expected adverse events based on real world and open-source data
- Inclusion of regulatory compliance reporting requirements

CTIS CTRM Domain Solutions

- Clinical Trial Planning & Management
- Data Integration, and Exchange
- Patient Enrollment and Retention
- Drug monitoring and reporting
- Document Management
- Regulatory Reporting & Analytics
- Site Selection & Investigator Management
- Adverse Event Reporting
- Data Governance, Security & Compliance

Protocol Development, LOI, e-Submissions, data extraction, Doc. Collaboration, Document Review workflow, Investigator and Site Management

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