CluePoints

Site Profile & Oversight Tool (SPOT)

SPOT is a site profile and oversight tool that enables adaptive site monitoring so teams can pinpoint anomalies and translate insights into actionable strategies. Clinical trials involve various stages, and as they progress, it becomes increasingly important to monitor them consistently and accurately to ensure the reliability of the data collected.

SPOT equips Sponsors and CROs with data-driven site monitoring for tasks that help surpass human capabilities. Discover how SPOT provides solutions to these common challenges.

CHALLENGE	CAPABILITY
Inefficient Monitoring Visit Planning	**Enable dynamic adjustment of site visits** based on anticipated site workload and central data review risk.
Time-Consuming Site Performance Evaluation	**Eliminate labor-intensive manual evaluations,** enabling study managers and CRAs to assess performance and take prompt corrective actions.
Unoptimized Site Recruitment Strategies	**Leverage valuable insights on site performance** to identify highly successful studies and replicate effective site selection and recruitment approaches.
Rigid Source Data Verification & Source Data Review (SDR) Strategies	**Tailor fit-for-purpose verification and review strategies** to meet unique study needs and risks, optimizing resource allocation and activities.
Inability to Evaluate CRAs Performance	**Assess CRA effectiveness across studies** with real-time site data, enabling more informed decisions on CRA assignments and training.

Reduce Site
Visits & Costs

Streamline System
Integration

Optimize
Cross-Study Analysis

Plan Future Study
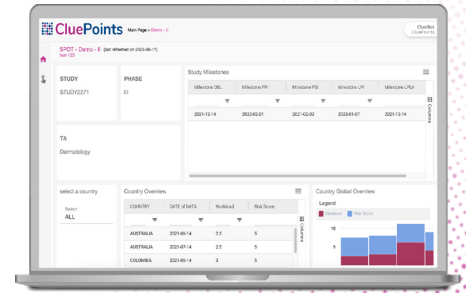Site Selection

Guide Critical
Thinking

Scan to Connect with an Expert
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Unlocking SPOT's Full Potential

SPOT delivers long-term consistency and control in site monitoring across various resourcing models and therapeutic areas. CluePoints guides you through every step of implementation with expert training and validation support, as well as custom dashboards, templates, and data transformation capabilities for improved oversight and efficiency. **While the benefits are extensive, here are our top five.**

1
DECREASE SITE VISITS TO INCREASE COST SAVINGS AND EFFICIENCY

2
BETTER PREPARE FOR SITE VISITS AND INTEGRATE MULTIPLE SYSTEMS

3
OPTIMIZE RESOURCE ALLOCATION WITH CROSS-STUDY ANALYSES

4
GUIDE CRITICAL THINKING FOR SITE MONITORING VISITS

5
MAKE INFORMED DECISIONS REGARDING FUTURE SITE SELECTION

See More Success with Central Statistical Monitoring

CluePoints' cloud-based solutions are driven by Central Statistical Monitoring, which leverages AI through advanced statistics and machine learning to interrogate clinical and operational data and illuminate data outliers and anomalies in real time. At the heart of our efforts are data quality and compliance monitoring. Sponsors and CROs can improve their ability to evaluate the performance of clinical trials and adjust site visitation plans more effectively and efficiently to consider risk and resource workloads more accurately.

DATA-DRIVEN SITE SCORING

USER-TRIGGERED ACTIONS

RISK MITIGATION

AUTOMATED ACTIONS

DATA VISUALIZATION

DATA INTEGRATION

ACCOUNTABILITY

ACCESSIBILITY FOR ALL

STUDY-LEVEL CONFIGURATIONS

FLEXIBLE & TECH-AGNOSTIC

Scan to Schedule a Personalized Demo
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