cliantha®
research

Clinical trials are evolving. Monitoring should too.

Because risks don't wait.

Because data doesn't lie.

Because decisions can't be delayed.

Cliantha.com

INDIA | USA | CANADA

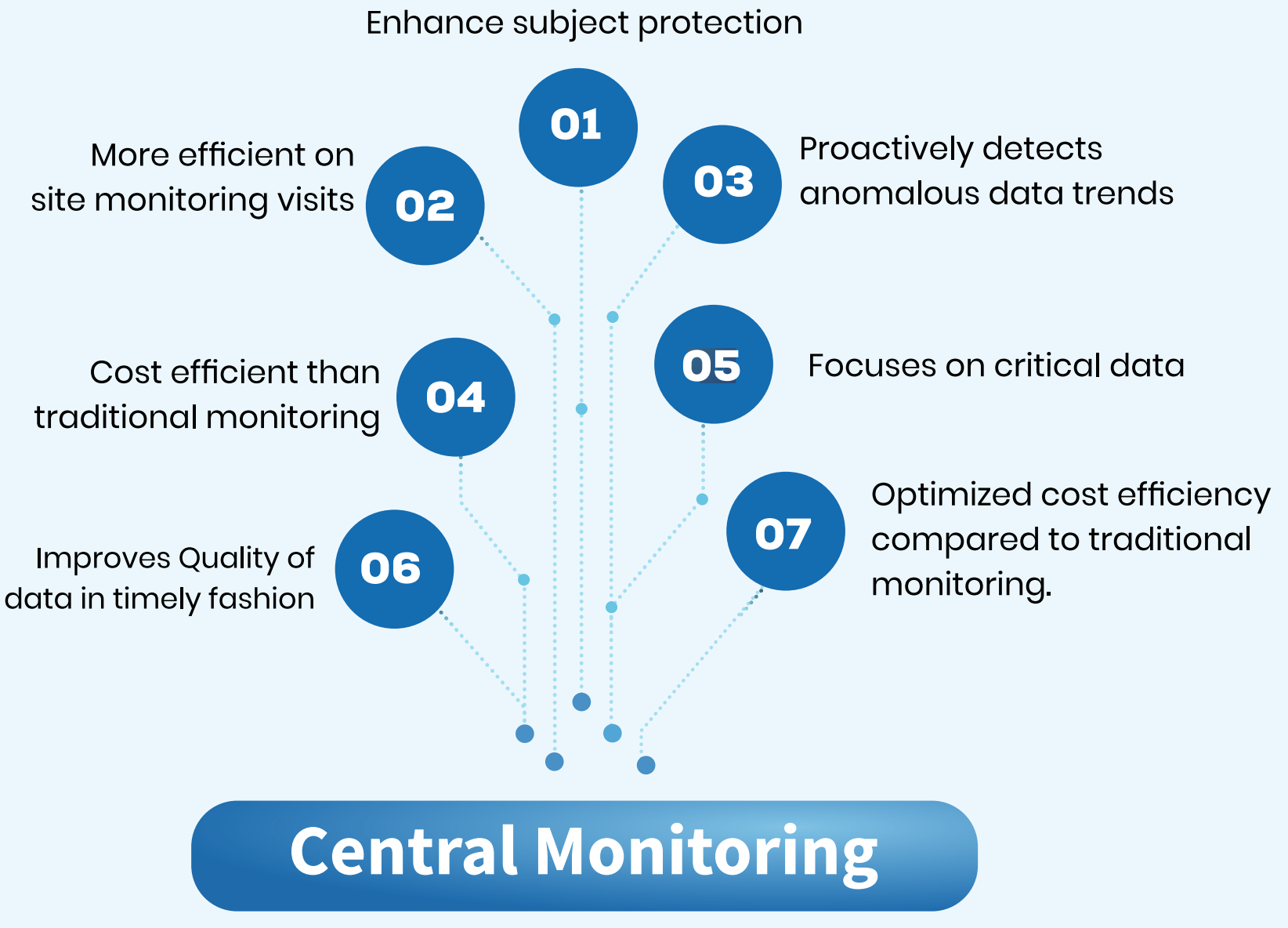
Clinical trials generate vast, complex, multi-site data.
But critical signals often remain hidden.

From site-based checks ⟶ Centralized intelligence

Central Monitoring

Central monitoring

Enhancing Clinical Trial Oversight with Data-Driven Precision!



At Cliantha
Expertise that goes beyond monitoring

Proactive

Data-driven

Insight-led

Our experience in Central Monitoring spans:

- Proactive risk identification and signal detection
- Data trend analysis and targeted oversight
- Cross-functional collaboration for faster issue resolution
- Driving inspection readiness through robust documentation

Aligned with regulations.
Driven by quality.

ICH E6 (R2)
Risk-Based Monitoring

Emerging ICH E6 (R3)
expectations

Focus on:

- Data Integrity
- Quality Risk Management
- Inspection Readiness

Why Cliantha?

Improved
Data Quality

Faster Issue
Resolution

Reduced
Site Burden

Enhanced
Trial Efficiency

Committed to delivering excellence through proactive monitoring, regulatory compliance and continuous quality improvement.

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