Vestigo® by McCreadieGroup


# General Information About Vestigo®

## What is Vestigo®?

Vestigo by McCreadie Group is a comprehensive software suite specifically designed to streamline protocol management across the entire Investigational Product (IP) supply chain. It encompasses a range of software solutions that significantly improve protocol management processes for research pharmacy sites, sponsors, CROs, and Interactive Response Technology (IRT) organizations.

## How does Vestigo® contribute to efficient protocol management?

Vestigo is a robust platform that allows research sites and institutions to effectively manage up to thousands of protocols and millions of transactions within the IP supply chain. It is designed to streamline and enhance the efficiency of protocol management processes, leading to improved workflows and better resource allocation.

## How does Vestigo® support key stakeholders in the Investigational Product (IP) supply chain?

Vestigo and its suite of software solutions support key players such as research pharmacy sites, sponsors, CROs, and IRT organizations through data transparency and integrity, and data standardization. By enhancing critical processes and facilitating seamless collaboration among these stakeholders, Vestigo ensures a smooth and efficient clinical trial management process across the IP supply chain.

## What are the key features and capabilities of Vestigo®?

- **Drug Accountability and Dispensing:** Elevates drug accountability and dispensing by guiding users through receipt, dispensing, return, and destruction of investigational products. Generates accurate Drug Accountability Record Forms (DARFs) and offers customization for labels.
- **Inventory Management:** Features push/pull system for alerts and reports, ensuring adequate supply levels, proper handling of products, and temperature excursion documentation.
- **Billing and Workload Management:** Enables creation of protocol-specific budgets, real-time charge recording, customized invoices, and automates time-based workload recording for clinical trials.
- **Protocol Management:** Facilitates checklist creation, task assignments, data standardization, and tracking of key protocol contacts.
- **Monitor Visits and Audits:** Centralized monitor database, integrated calendar for scheduling visits, and robust documentation with audit trails.
- **Document Management:** Upload, create, and organize essential documents, maintain temperature logs, and provide a long-term storage solution for electronic documents.

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## Contact Us

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