

goBalto Activate

Driving Site Activation

Accelerating site activation can jump-start a clinical trial, but with the hours and weeks spent manually tracking tasks and documents without visibility into where the bottlenecks lie, how can you make that happen?

Improve Operational Performance

Study teams are under intense pressure to speed clinical trials and restrain costs, but inefficiencies tied to complicated protocols, globalization and paper-based methods have stalled these efforts.

goBalto Activate improves operational efficiencies with real-time monitoring of items on the critical path to ensure key milestones are met.

With goBalto Activate, you can:

- Leverage 70+ standardized country workflows for quick site activation.
- Track any activity, document, submission, or ad-hoc documents.
- Drive optimization with role-based assignments and dependency triggers.

Streamlined Document Management

The “old way” of using email, spreadsheets, and e-rooms to track documents and activities associated with activating study sites does not work for today’s clinical trials. Study teams need real-time access to documents and automated, compliant workflows.

goBalto Activate lets you:

- Know you’re working with the right file with document lineage.
- Select documents for IP release reviews or submission packages.
- Proactively request new documents prior to expiry with alerts.



“goBalto Activate’s robust, industry-proven library of country-specific workflows and management-based approach to site activation is central to our efforts to build out our infrastructure of ‘best of breed’ applications in our eClinical suite. Using a workflow-based approach to study startup, critical indicators of quality can be assessed on an ongoing basis so that corrective actions, if needed, can be made earlier.”

MIKE TOWNLEY
CIO
CLINIPACE



Actionable Data, Readily Available

When a clinical trial does not unfold as planned, it may be heading for rescue. The key to avoiding study rescue is proper identification of red flags signaling a study may be veering off course.

- Real-time study progress reporting drives transparency and highlights opportunities ripe for optimization

goBalto Activate streamlines and automates site activation, cutting cycle time by over 30%.

goBalto Activate provides:

- Real-time study progress reporting drives transparency and highlights opportunities ripe for optimization.
- Critical path tracking reports show which activities delay key milestones.
- Alerts notify study team members when pre-requisite work is completed.

Drive Collaboration and Deliver Compliance

Sponsors often have multiple studies running concurrently with multiple CROs, complicating oversight. Having open and transparent communications across all your trial partners is critical, but this is difficult when you're tied to slow, manual processes.

Foster open communication and trust with goBalto Activate by:

- Providing a centralized view of all assigned study team members.
- Exchanging documents, notes, completed activities, and monitor partners status with audit trail reporting.
- Ensuring compliance with organizational SOPs and country-specific regulations.

Oracle Health Sciences

Oracle Health Sciences breaks down barriers and opens new pathways to unify people and processes, helping to bring new drugs to market faster. As the #1 provider of eClinical solutions (Everest Group, 2017), and powered by the #1 data management technology in the world (Gartner, 2018), Oracle Health Sciences is trusted by 29 of the top 30 pharma, 10 of the top 10 biotech, and 10 of the top 10 CROs for clinical trial and safety management around the globe.





AUTOMATE WORKFLOW

Intelligent workflows drive study teams to complete and track specific documents and tasks required for any site, country, or study based on regulatory requirements and standard operating procedures.

LEARN MORE

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