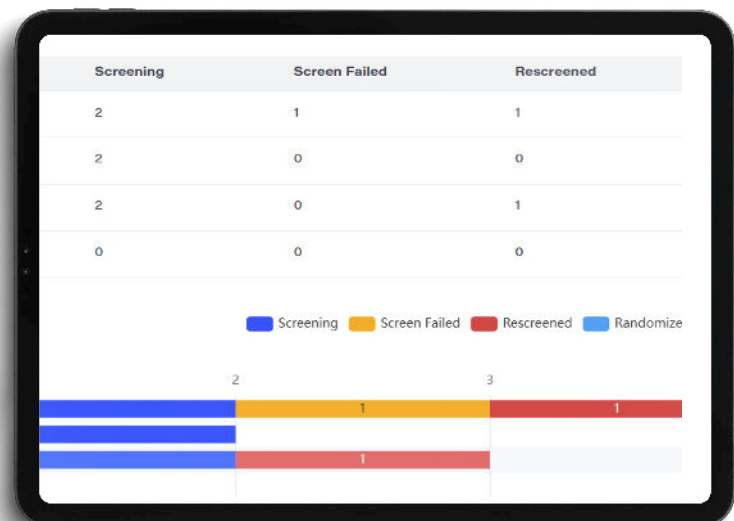


Introducing ClinPhone Pro

ClinPhone Pro supports Study Managers with:

- ▶ **Flexibility** - adapt instantly with self-service tools, keeping scope changes minimal.
- ▶ **Efficiency** - track study progress with real-time visibility - from system access ahead of Site Initiation Visits, to progress against enrolment targets, to participant data change history.
- ▶ **Easy from start to end** - reduce your specification review and UAT time with early system demos and pre-validated libraries tailored to study design, agreed standards, Therapeutic Area and Phase.
- ▶ **Site-friendly** - intuitive design boosts efficiency and accuracy, with calendar views of visits, guided visit registration and tooltips. Past study notifications can be re-sent at any time, supporting site audits.
- ▶ **First-time data quality** - built-in validation ensures first-time accuracy. Comprehensive reports provide full traceability of all data changes.
- ▶ **Reports for your every need** - instant access to data you need, from dashboards showing high-level status to downloadable reports showing participant visits and full audit trails.



Book a demo of this future-proof RTSM solution

Introducing ClinPhone Pro

30+ years of IRT expertise - delivering secure, reliable, and precise randomization and trial supply management.

A future-proof IRT solution enabled by clever architecture and a trusted industry cloud for fast, scalable, and resilient development.

Confidence that the right participant gets the right treatment at the right time - every time, plus:

- ▶ Rapid study build
- ▶ Effortless set-up with study demos from day 1
- ▶ Configurable design tailored to your study, Therapeutic Area, and Phase
- ▶ 1-click study design using pre-validated templates
- ▶ Self-service for instant study design changes
- ▶ Intuitive, user-friendly interface
- ▶ On-demand reporting from status and dashboards to audit trails
- ▶ Seamless system integration with supply, EDC and other systems

Expert support - from RTSM specialists, to an 24/7 in-house helpdesk.

Committed to sustainability and on-time data submission, with 500+ regulatory approvals.

Your trusted ally - from day one to database lock.



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