Trial wisely

Our IRT Reduces Change Orders

Make mid-study changes quickly and easily. Pivot with protocol amendments and adapt your trials with fewer change orders.



Consultative Approach System builds that meet your needs



eClinical Expertise Comprehensive, high-quality systems



Flexible Platform Configurable systems to manage the unexpected





LEARN MORE Visit suvoda.com

to learn about our solutions

The following system updates can be made through our IRT interface:

STUDY AND SITE

- · Activate and deactivate sites
- Add sites
- · Modify site or investigator data
- Modify study enrollment and screening limits
- Modify country enrollment and screening limits
- Modify site enrollment and screening limits
- Add and modify satellite sites associated with a central pharmacy*
- Add and modify study doses*
- Add and modify cohorts*
- Create, modify, save, and share ad-hoc reports

ADMINISTRATIVE

- Modify users' access and role
- Approve user access
- Add and modify notification subscriptions
- Add and modify user roles
- Modify system role permissions
- · Correct subject or drug data

DRUG MANAGEMENT

- Add depots and add lots
- Release drug incrementally, as required
- Modify depot distribution
- Add and modify supply strategies
- Modify supply strategy association with sites
- Modify supply strategy advanced rules for enrollment visit(s)*
- Add and modify dose modification rules*
- Modify lot expiration dates
- Add and modify drug label groups
- Add and modify rules for temperature excursions*
- Add and modify drug destruction facilities*
- Configure specific sites, depots, and/or countries to receive certain drugs from specific depots*
- Release or withhold drug for use by country and/or globally