SUVODA®
Trial wisely

Your oncology trials, simplified

Our technology promises a simpler oncology trial for your study team, so that your patients are truly at the center of everything you do.

Intuitive IRT Interface
Gracefully navigate your supply chain complexities

Flexible System
Manage your inevitable mid-study changes with ease

Unified on the Suvoda Platform
A single data model and streamlined workflows across eConsent, IRT, eCOA, and ePatient

Powerful IRT functionalities for unpredictable oncology trials

DYNAMIC VISIT SCHEDULE AND CYCLE EXPANSION

- Automate cycle expansion and variable visit schedules
- Change patient visit schedules on the fly directly through the IRT interface

EARLY PHASE STUDIES: DOSE AND DISPENSATION MANAGEMENT

- Add and change dose and dispensation configurations mid-study
- Indicate kit quantity allotted at each dispensing visit

COMPARATOR AND ADJUVANT THERAPY MANAGEMENT

- Account for different sourcing of medications by site or country
- Add doses and configure dispensations for comparators mid-study

ADVANCED COHORT MANAGEMENT

- Easily control open cohorts and enrollment limits
- Add new cohorts and dose levels mid-study without programming changes

DISEASE TYPE MANAGEMENT FOR BASKET STUDIES

- Add new disease and tumor types in real-time within the IRT interface
- Open and close disease types for enrollment and set enrollment limits

THERAPEUTIC EXPERTISE

100%
Services teams with oncology experience

43%
Studies in oncology

52%
Oncology studies that are early phase

112k+
Oncology patients worldwide

GLOBAL EXPERTISE

1,800+
Trials supported

500+
Clients

100k+
Sites using Suvoda

95
Countries supported

100%
Client milestones achieved on time

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