ꟿUVODA

Trial wisely

Evolve Your Clinical Trials with Direct-to-Patient Logistics

Our field-tested IRT functionality allows you to strengthen patient engagement and navigate study complexity with ease.

End-to-end Visibility
Robust supply chain management tools that uphold blind and patient confidentiality

Fast, Flexible Builds
Best-of-breed functionality and integrations that streamline drug and patient logistics

Multi-therapeutic Experience
Well-versed in offering direct-to-patient functionality across therapeutic areas

What our Direct-to-Patient functionality gives you

RIGOROUS CONTROL FOR HYBRID AND DECENTRALIZED TRIALS

- Toggle direct-to-patient functionality for your IRT system on/off at the site, country, and/or patient level as needed.

- Configure site-to-patient, depot-to-patient, and central pharmacy-to-patient shipments, or a combination of these depending on country or region-specific requirements.

INTUITIVE DRUG SUPPLY MANAGEMENT TOOLS

- Ship drug to patients manually or automatically based on specific dates built into the visit schedule.

- Drug supply managers and site users can request additional shipments as needed.

- Site users and in-home staff can track receipt of shipments to patients.

INTEGRAL IRT FEATURES THAT ENHANCE LOGISTICS

- Build ad-hoc reports and access pre-built, real-time reporting to maintain compliance and identify data trends.

- Integrate with EDC systems to efficiently manage clinical data.

- Manage open-label studies and blinded studies with built-in blinding protections for common direct-to-patient scenarios.