SUVODA®

Trial wisely

IRT + eConsent: A Simplified Process

A streamlined approach to realizing the benefits of eConsent

- One trusted, consultative project team
- Single UAT for both IRT and eConsent
- No impact to IRT project timelines

SUVODA®

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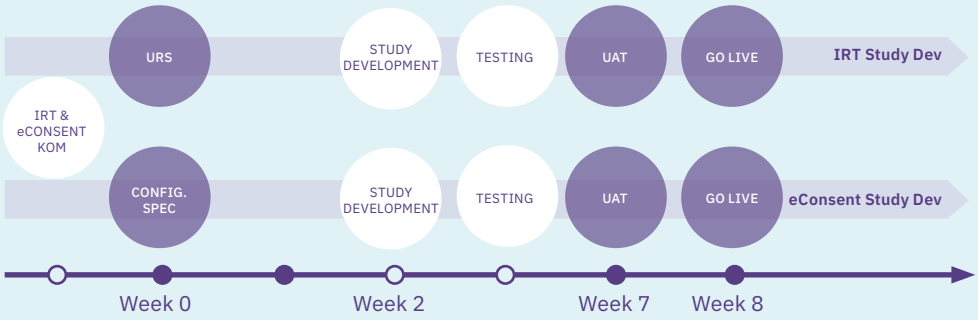
to learn about our solutions

IRT + eCONSENT: NO IMPACT TO IMPLEMENTATION TIMELINES

Suvoda eConsent, a flexible solution designed to adapt to requirements across the globe, can be added to the scope of your IRT project with no impact to implementation timelines. IRT + eConsent builds follow the same implementation processes and have the same project milestones, meetings, and deliverables.



STREAMLINED PROCESS FOR EFFICIENCY

+ **One Team:** A single delivery team and Project Manager leading you to success
+ **Requirements Gathering:** IRT user requirement specifications (URS) are gathered alongside the eConsent configuration specification
+ **Startup:** Joint IRT + eConsent kick-off meeting
+ **UAT:** Perform your User Acceptance Testing on Suvoda's entire platform, seeing patient scenarios lead you from eConsent to IRT

Sponsors and CROs can go-live with the IRBs, ECs, consent forms, and languages that are available at that time.

Post go-live, consent forms, languages, and IRBs are added just-in-time by the Suvoda project team, no change orders required. Mid-study protocol amendments resulting in new consent forms are seamlessly pushed to specified sites.

SERVICE EXCELLENCE

Sponsors and CROs can lean on the trusted, consultative Suvoda project team, which includes an eConsent specialist, to be intimately aware of the nuances of your trial and protocol.

Following go-live, the same project team will support the study team throughout the lifespan of the trial. With a median resolution time of less than 19 minutes for high priority help desk tickets and 100% of our services team having complex clinical trial experience, sponsors and CROs can expect a seamless and connected experience throughout their trial.