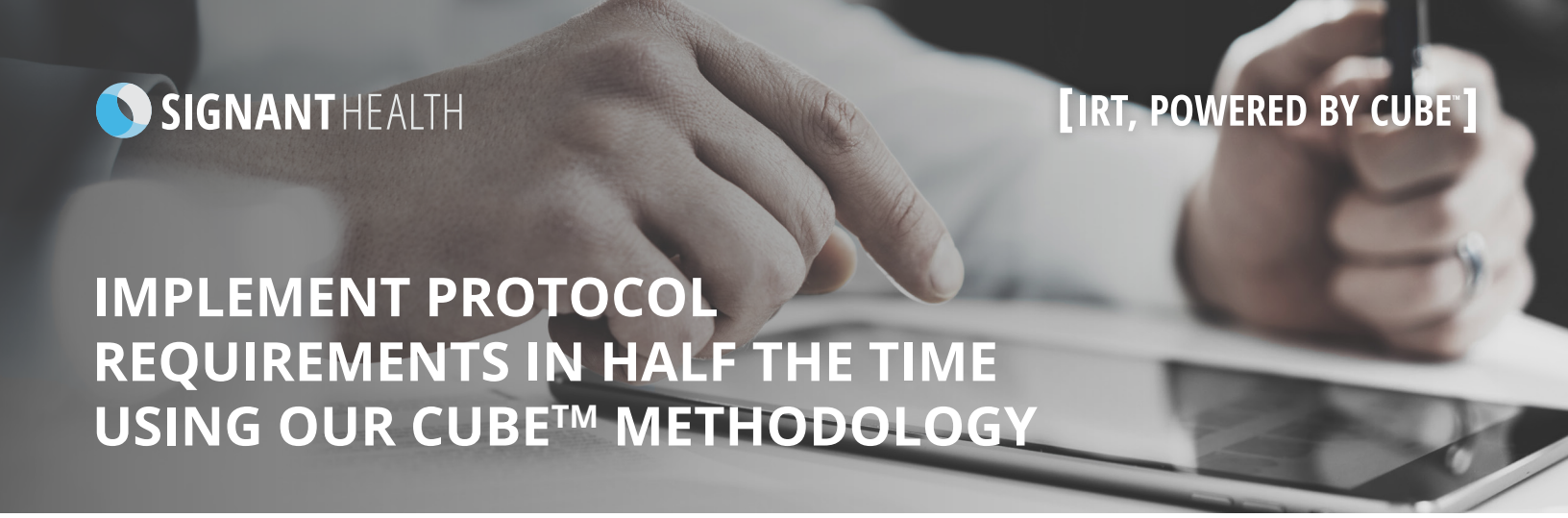


SIGNANT HEALTH

[IRT, POWERED BY CUBE™]

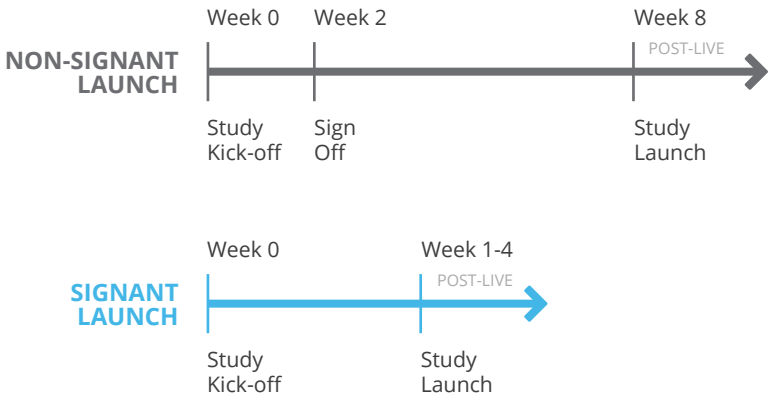
IMPLEMENT PROTOCOL REQUIREMENTS IN HALF THE TIME USING OUR CUBE™ METHODOLOGY

Signant Health's IRT is a quick-deploy solution for managing randomization and clinical supplies, trusted by leading life science companies for more than 20 years. Its proprietary deployment framework, CUBE™, which stands for Configuration by User Based Experience, is a game-changing delivery method. It offers the industry's fastest system setup process while ensuring even your most complex requirements are ready and to your liking at go-live. The system can be accessed at any time via the web or mobile app, providing additional flexibility to end users.

The Signant IRT system can be deployed with CUBE™ on any protocol, including those with complex randomization schemes, advanced dosing schedules and custom requirements. Our typical deployment timeline is 1 to 4 weeks from the initial kick-off meeting.

The CUBE™ pre-validated configuration modules and agile implementation methodology allow for faster, more accurate deployment and maintenance of your IRT system.

2X FASTER IRT LAUNCH WHEN DEPLOYED WITH CUBE™



2X FASTER DELIVERY AND 50% FASTER MID-TRIAL CHANGES

We can deploy your IRT system in 1 to 4 weeks from the initial kick-off meeting. When protocol amendments happen mid-study, we use the same process to deploy system updates quickly.

50% LESS STUDY TEAM EFFORT REQUIRED AT STUDY STARTUP

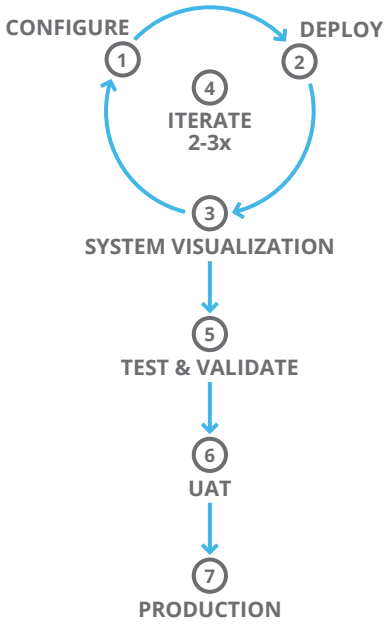
We know your time is especially valuable during the study startup phase and have therefore developed configurable checklists to save you hours of design calls and allow you to skip review of source documents.

100% SATISFACTION THAT YOUR REQUIREMENTS ALIGN

You can visualize the system as it is being built for your protocol and we make updates in real-time, eliminating risk and ensuring the system we launch matches your requirements perfectly.

HOW CUBE™ WORKS TO LAUNCH YOUR IRT SYSTEM

The CUBE™ methodology is an iterative process that leverages a comprehensive library of pre-validated functionality and custom features built on-demand to meet the needs and complexity of any clinical study.



OUR ITERATIVE CONFIGURATION EXPERIENCE ENSURES YOUR IRT SYSTEM MEETS EXPECTATIONS

1. **CONFIGURE**
   We activate the features you need by switching each one on/off in our extensive pre-validated code library
2. **DEPLOY**
   You are given access to the study-specific system
3. **SYSTEM VISUALIZATION**
   You can visualize the system and request configuration changes
4. **ITERATE**
   This cycle is repeated 2 to 3 times until you approve the final configuration
5. **TEST & VALIDATE**
   We test and validate any custom elements against the specifications
6. **UAT**
   You conduct a final UAT of the system prior to launch
7. **PRODUCTION**
   Your system is ready for launch

COMPATIBLE WITH EDCS, EPROS, CTMSS, OTHER DATA SYSTEMS AND SMARTSUPPLIES®

We ensure that integrations between Signant's IRT and your study's other data system are up and running for launch. We are the only provider to offer an out-of-the-box integration between IRT and SmartSupplies, a solution that forecasts, labels, distributes, manages and reconciles clinical supplies. Our clients use these solutions together to avoid stock outs, reduce overage and prevent unnecessary packing runs.

THINK AGAIN, PAPER IS NOT THE ONLY LIGHTWEIGHT SOLUTION

Are you using spreadsheets because traditional IRT systems are costly and slow to setup? With the power of CUBE™, Signant's IRT can be quickly and easily configured to function as a lightweight system to support small Phase I studies, investigator-initiated trials and more.

24/7 SUPPORT

At Signant, we recognize that leading edge technology is only part of the success criteria for IRT deployment. We provide industry- leading support via a dedicated project team and 24/7 helpdesk to ensure your success from system build to study close.

WHO IS SIGNANT HEALTH?

The best technology succeeds in the background. Signant Health provides solutions that simplify every step of the patient journey to make it easier for people to participate in, and for sites and study teams to run, clinical trials. Signant unites eCOA, eConsent, Patient Engagement, IRT, Clinical Supplies and Endpoint Quality into the industry's most comprehensive patient-centric suite – an evolution built on more than 20 years of proven clinical research technology. Our intense focus on the patient experience, deep therapeutic area expertise and global operational scale enable hundreds of sponsors and CROs (including all Top 20 pharma) to extend the reach of drug development, expand patient opportunities and improve data quality – helping them bring life-changing therapies to our families and communities around the world. Take a significant step toward patient-centricity at signanthealth.com.

CRF Health and Bracket are now Signant Health.

SIGNANT HEALTH