Fact Sheet

# Orchestrated Study Start-Up with the IQVIA Investigator Site Portal

*Reduce cycle time, speed site activation, and keep sites engaged with robust workflows and compliant eTMF filings*

Orchestrated Study Start-Up with the IQVIA Investigator Site Portal reduces activation timelines and eliminates site frustrations by automating processes and providing complete oversight to study teams. Complex processes for start-up and beyond are streamlined through feature-rich technology that sites and sponsors are eager to use.

## Activate studies, countries, and sites More quickly

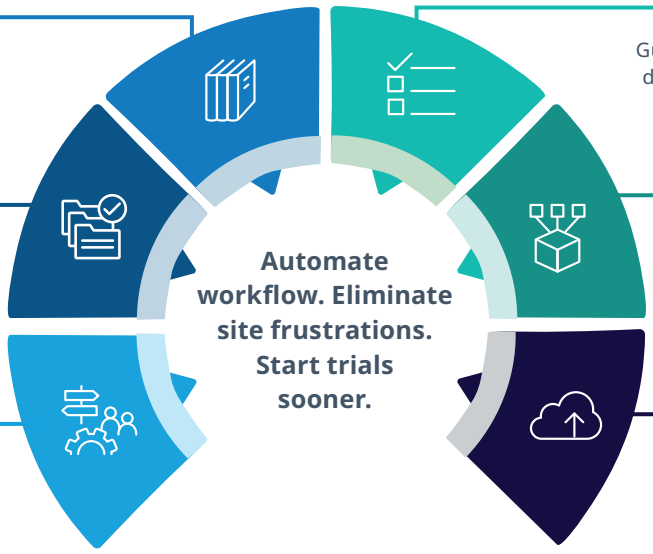
Orchestrated Study Start-Up accelerates time to activation, enables site-sponsor collaboration, and empowers study teams to make better informed decisions. It provides highly configurable study, country, and site workflows while ensuring all documents destined for the electronic trial master file (eTMF) are sent promptly upon finalization for quality control and contemporaneous filing, with ongoing management of key performance indicators (KPIs) for continuous process improvement.

## Deploy the tech that sites already know and love

For more than a decade, the IQVIA Investigator Site Portal (formerly DrugDev) has been the site-facing technology that sites love to use. More than 150,000 investigators and site staff sign in at an average rate of 96% — and it's so simple to use that no additional training is ever required.

Now with advanced features for site, country, and study activation, the Investigator Site Portal is the first choice for sponsors that need to orchestrate study start-up across their portfolios.

**Key features of advanced site activation**




**CTMS Orchestration**

Automate user activities and track start-up data in your system of record with bi-directional CTMS orchestration

**Global Catalog**

Create consistent start-up processes and metrics collection across all trials and enable study teams to set up even large studies in a day

**Study configuration**

Select templates from the Global Catalog to configure start-up activities for varying study requirements, therapeutic areas, or countries

**Document Checklists**

Guide study team and site users through document collection and approval using visual actions icons, task-based filters, and notifications

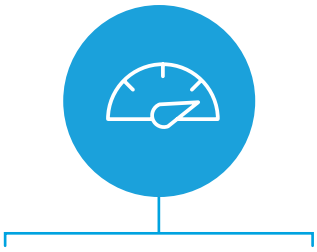
**Submission milestones**

Track groups of major requirements/ approval activities such as regionspecific regulatory submissions in a dashboard

**ETMF Orchestration**

Transfer approved country and site documents to eTMF systems including required metadata values

**Benefits**



### Faster study start-up

Streamline feasibility, site selection, ethics committee approvals, and document collection in a single platform

### Closer collaborations

Achieve greater alignment across study partners with a seamless exchange of trial information that drives unmatched site satisfaction

### Better decisions

Bring all the data together and get end-to-end oversight in an intelligent analytics platform

## A complete solution from start-up to close-out

The IQVIA Investigator Site Portal is a one-stop shop for study start-up, conduct, and close-out, proven to provide significant value to sites and sponsors in thousands of trials.

**Site identification and selection**
Expedite site selection decisions with built-in workflows that automate CDA acceptance and increase survey responses.

**Site engagement**
Improve patient enrollment and retention rates, protocol adherence, and data quality with trial communications, site-first tools, and automated trackers.

**Activation**
Get to first patient/first visit faster and eliminate site frustrations by automating processes and enabling oversight for study teams.

**Safety notifications**
Disseminate and track safety notifications with complete transparency, without overwhelming investigators and site staff.

**Learning Management**
Ensure site compliance and satisfaction through automated assignments, cross-trial credits, and comprehensive reporting.

**Document library**
Give sites self-service access to all the information they need to be successful, with powerful sharing, searching, and tracking for efficient document exchange and team collaboration.

# Orchestrate Outcomes

IQVIA TECHNOLOGIES

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