FACT SHEET

IQVIA Feasibility

A data-first approach optimizing feasibility and site selection

Transform feasibility using a digital approach

Employing a digital, data-first strategy, IQVIA Feasibility brings together survey data, confidential disclosure agreements (CDAs) and site intelligence data, enabling sponsors to efficiently collect vital information while improving the site experience. Powered by IQVIA Connected Intelligence™, the system uses purpose-built, automated technology and advanced analytics to leverage the information so you can make the right decisions regarding trial strategy and planning, protocol design, and country and site selection.

EASE SITE AND SPONSOR BURDEN

This cloud-based solution improves the sponsor, site and investigator experience, while providing increased clarity and better oversight.

HEIGHTEN YOUR PROBABILITY OF SUCCESS

- Standardize and streamline processes
- Centralize and automate data and document exchange
- Optimize site-sponsor communications

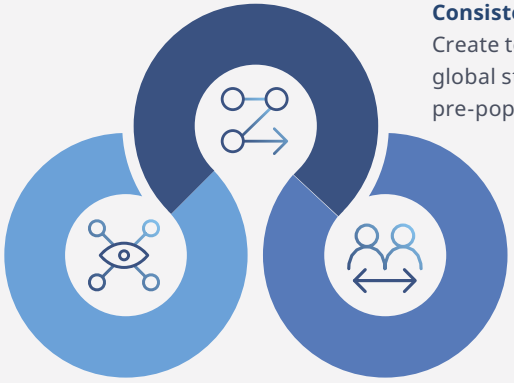
- Simplify CDA and survey development
- Shorten critical process timelines

UNLOCK DATA. AUTOMATE WORKFLOWS. ELIMINATE SITE FRUSTRATIONS.

SURVEY DESIGN AND REVIEW	SITE OUTREACH	CDA WORKFLOW	SURVEY RESPONSE	ANALYSIS AND SELECTION	DASHBOARD AND REPORTING
Standardize and re-use survey questions and leverage advanced survey review capabilities to improve collaboration.	Better orchestrate outreach and increase transparency for efficient follow-up.	Automate workflow with eCDAs, in-tool negotiation, document exchange and tracking.	Survey pre-population speeds site response.	Assess investigator suitability with multi-sourced information and dynamic scoring; capture site selection decisions.	Export real-time reports at site, country or study level for further analysis.

Purpose-built to optimize your clinical trials

Centralized, standardized feasibility assessments
Single solution for end-to-end processes, bringing together surveys, CDAs and intelligence data

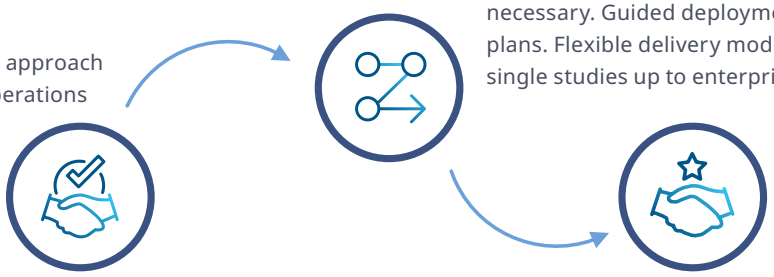


Consistent processes with automated workflows
Create templates which can be used to drive global standardization and allow for survey pre-population and eCDA automation

Deepen collaboration and operational effectiveness
Optimize operations across countries, regions and global teams with a bespoke feasibility solution

Keep your studies on track

Expertise
Consultative, partnered approach optimizing feasibility operations across people, process, technology and data.



Simple implementation
No technical requirements or integrations necessary. Guided deployment and adoption plans. Flexible delivery models. Available for single studies up to enterprise partnerships.

Become sponsor of choice
Foster site relationships and ease site burden by simplifying delegation, streamlining the CDA process and leveraging existing site response data via survey pre-population.

Added value for all stakeholders

Centralize internal team efforts
One survey can be managed across regions to enable local feasibility/site ID collaboration and centralize data output

Streamline CDA negotiation
Via eCDAs and document CDAs, in-system document exchange, version control and document management

Effective and efficient site follow up
Automatic tracking of CDA and survey. All stakeholders have transparency with visible follow up history

Increase survey response
Pre-population of survey responses from previous surveys in an easy to use interface

Reduce site selection timelines
Accelerate cycle times of survey creation, invitation process, CDA negotiation and survey completion, leading to site selection

Improve investigator experience
Direct sponsor-site interaction, simplified CDA negotiation, survey completion and survey management across sponsors

Expedite intelligent site selection
Dynamic scoring and ranking enables faster, more protocol-focused site selection including documentation of rationale for future intelligence

For more information on how IQVIA Feasibility can help improve your clinical trials' probability of success, please contact us today to speak with an expert: orchestrateyourtrials@iqvia.com

IQVIA TECHNOLOGIES

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