

IQVIA Planning Suite

Design your clinical trials with optimized precision

Overcome study design challenges

Despite heavy investment, many clinical trials either fail or require extended timelines to complete. Challenges with study design and feasibility, access to robust data sources, country and site selection and budgeting often lead to site under-performance and dissatisfaction, poor patient enrollment and millions of dollars in lost revenues per day.

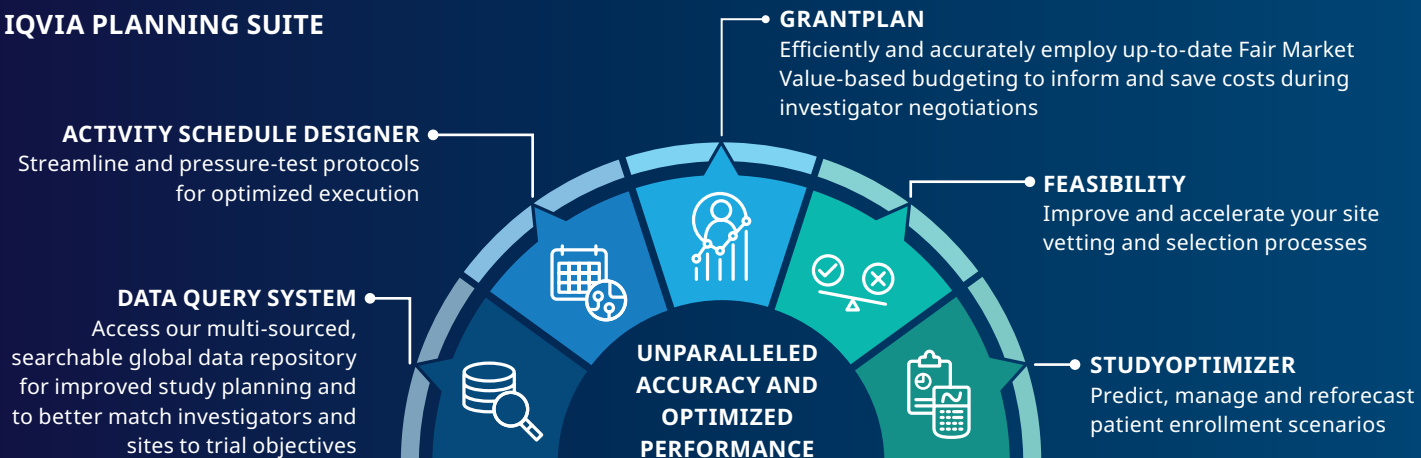


OPTIMIZE ALL ASPECTS OF CLINICAL TRIAL DESIGN AND PLANNING

The IQVIA Planning Suite employs purpose-built technologies, the world's largest, most up-to-date repositories of high quality, multi-sourced, clean and curated data along with decades of clinical trial experience to help life sciences companies and service

providers accurately predict operational execution and budgeting to keep studies on track. The suite consists of five SaaS solutions that help optimize all aspects of clinical trial design and planning across trial phases, therapeutic areas and world geographies. Forecast study timelines, costs and performance with greater precision to make better decisions and drive smarter, faster trials.

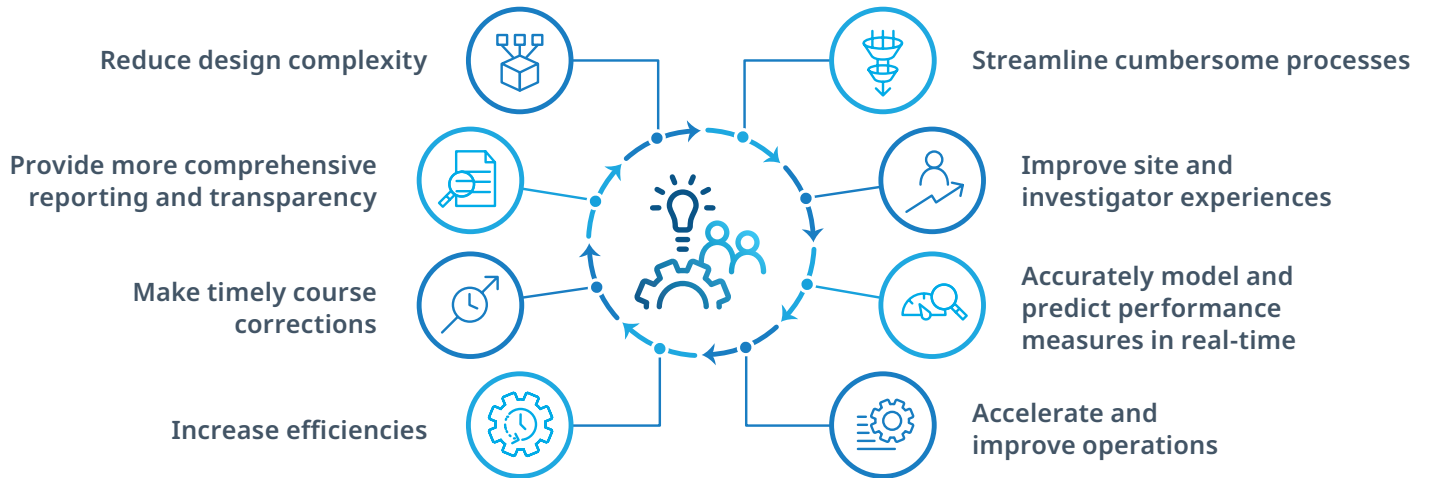
IQVIA PLANNING SUITE



Orchestrate better study outcomes right from the start with IQVIA Planning Suite and enjoy greater return on your investment in clinical development.

BETTER PLANNING & PERFORMANCE





Employing these intuitive, evidence-based tools allows users to further:



INTEGRATE WHERE IT MAKES SENSE

IQVIA's Planning Suite affords the flexibility to integrate where it makes sense for your organization — whether you're looking for an end-to-end SaaS solution, a single, best-in-class offering or tech-enabled service. The suite's orchestrated, proven, site-focused tools scale to meet client needs in over 100 countries and integrate with each other, third-party offerings and your existing IT environment.

BETTER STUDY EXECUTION RIGHT FROM THE START

- More accurate budgeting**  **76%** of global trials in clinicaltrial.gov use GrantPlan
- Fewer protocol amendments**  Up to **40%** reduction in protocol amendments and as much as **\$12K** lower per patient cost
- Reduced patient burden**  As much as **24%** reduction in patient burden scores
- Accelerated site selection**  Site selection completion accelerated by up to **30** days; faster survey completion by as much as **13** days

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