DrugDev
an IQVIA company

WHAT YOU GET WHEN YOU CHOOSE DRUGDEV SPARK

*The following data has been collected from webinars, case studies and other projects with DrugDev Spark customers.

SITE ACTIVATION

40%
Sponsors can see an average 40% reduction in contracting cycle time.

30%
Average time required to get all contracts completed can be reduced by up to 30% **per study**.

24 HOURS
"Using DrugDev Spark, I was done with the contracting process in 24 hours. It took four weeks for a similar trial with a different organization."
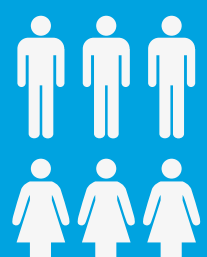— Nancy Baker | Site Director
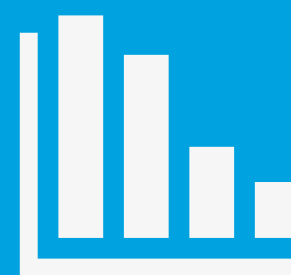
SITE ENGAGEMENT

50%
Reduce low-enrolling and non-enrolling sites by as much as half.

3x
Multiple sponsors have seen more patients enrolled by inspiring site performance.

Eliminate manual, error-prone, time-consuming tasks like browsing paper files, unwieldy online systems and seeking answers.

REDUCE MONITOR VISITS BY 15%

REDUCE PROTOCOL DEVIATIONS AND ENSURE CLEAN TRIAL DATA

SITE COMPLETE ALL CLOSE-OUTS VISITS IN AN AVERAGE OF 6-8 WEEKS

REDUCE TIME SPENT ON REPORTING AND UPDATES

SITE PAYMENTS

30%
Sponsors can save up to 30% in trial costs through VAT, cash flow optimization and FTE time savings.

INCREASED SITE SATISFACTION
By paying sites monthly instead of quarterly, sponsors can increase site satisfaction rates by as much as **80%**.

80%
AGGREGATE SPEND
Complete aggregate spend reports for global studies 80% faster.

> **DrugDev Spark is the best site technology on the planet. We contract with a variety of sponsors and CROs and when it comes to the financial lifecycle no one offers the level of detail that DrugDev does. That alone is enough to recommend, but the outstanding customer service that comes with it makes DrugDev the best in the industry.**
>
> *Sandy Zipp | Director of Study Accounting | Synexus USA*

eCONSENT

Enroll fewer patients to meet your same completion goals due to improved retention.

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Zero audit findings with eConsent.

Significant increase in patient satisfaction.

THE MAJORITY OF PATIENTS SURVEYED RATED ECONSENT EASY TO USE.

IMPROVE OVERALL INTEGRITY OF THE CONSENT PROCESS

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