

CASE STUDY: *STREAMLINING MULTISITE CLINICAL RESEARCH OPERATIONS*



CHALLENGE

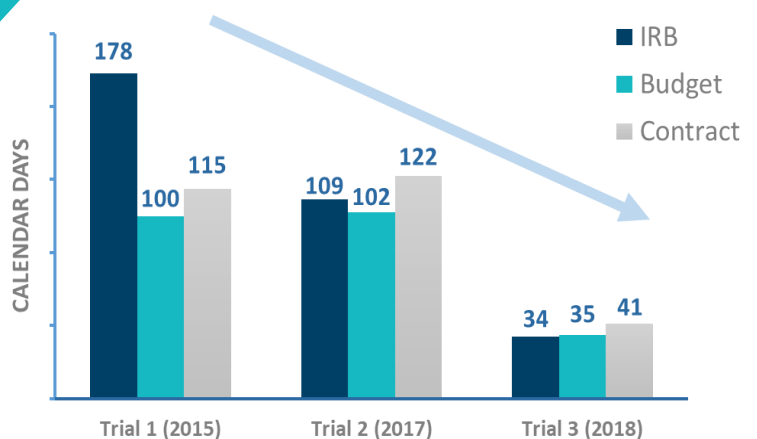
- PREMIER HEALTHCARE SYSTEM SOUGHT TO LOWER THE HURDLES FOR PHYSICIAN-INVESTIGATORS AND INDUSTRY SPONSORS TO CONDUCT MULTI-SITE CLINICAL TRIALS AND PROMOTE COLLABORATION.

SOLUTION

- DEVELOPED ENTERPRISE-WIDE CLINICAL TRIAL STUDY START-UP PROCESSES AND BEST PRACTICES TO DECREASE STUDY START-UP TIMELINES AND LIMIT TRADITIONAL ADMINISTRATIVE REDUNDANCIES.
- STANDARDIZED NETWORK CONTRACTUAL AGREEMENTS, REGULATORY AND BUDGETARY PROCESSES TO ACTIVATE MULTISITE STUDIES WITH 1 IRB SUBMISSION, 1 CONTRACT AND 1 COMMON STUDY BUDGET.
- PROACTIVELY EXECUTED MASTER AGREEMENTS WITH INDUSTRY PARTNERS AND INCREASED ADOPTION RATE OF CLINICAL RESEARCH TOOLS (I.E., SMARTIRB, ACTA, CTMS AND E-REG BINDERS).

IMPACT

- EXPANDED NETWORK COLLABORATION AND INCREASED CLINICAL TRIALS PORTFOLIO BY >250%.
- DECREASED STUDY START-UP ACTIVATION TIMELINES FOR INDUSTRY TRIALS BY 76% .
- INCREASED STUDY BUDGETS BY 46% COMPARED TO ORIGINAL SPONSOR BUDGETS.



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