

## The Right Fit For You.®



The experienced team at SDC understands that a clinical trial “rescue” raises unique challenges for the study team. With a consultative approach to rescue studies and scalable services to fit your needs, SDC is The Right Fit For You.®



### Rescue Study Services

- › Strategic Consultation for evaluating rescue options and service needs
- › Development and Implementation of efficient project transition timeline
- › Scalable Services ranging from consulting through full service CRO support

### Rescue Study Experience

- › Multi-study Rescue of full data services
- › Mid-study Transitions of Statistical Services
- › Gradual Transition of All Data Services across multiple programs
- › Assumed Responsibility for Existing EDC Database on global oncology study
- › Rescue Consultant for multiple programs



9 out of 10 Average Customer Satisfaction Score

98% On-time Delivery of Key Project Milestones

10+ Rescue Studies Supported in Past 2 Years

## The Right Fit For You.®



SDC is a specialized data services CRO providing scalable full-service clinical trial solutions via our diverse and complementary strategic partnerships. From protocol consulting to full-service clinical trial management and everything in between, SDC is The Right Fit For You.®

### Core Internal Services










- › **Biostatistics**
  - › Consulting Support
  - › PK/PD Analysis
  - › DMC/DSMB Statistics Services
  - › Submission-Ready Data Services
- › **Clinical Data Management**
  - › CDISC/CDASH Expertise
  - › Proactive Data Cleaning
- › **eClinical**
  - › Electronic Data Capture (EDC)
  - › Interactive Response Technology (IRT)

### Partnered Services

- › **Clinical Operations**
  - › Clinical Monitoring
  - › Project Management
  - › Medical Monitoring
  - › Site Management
- › **Regulatory Consulting**
- › **Medical Writing**
- › **DMC Assembly & Administration**
- › **Pharmacovigilance**

## Mini Case Study

› Two Pivotal Rescue Trials running in parallel

Study #1		
Item	Duration*	Timeline
eCRF Design, Development, and Approval	26 Days	
DMP Development and Approval	12 Days	
DVM Development and Approval	17 Days	
Release of EDC/IWRS into Production	39 Days	
Study #2		
Item	Duration*	Timeline
eCRF Design, Development, and Approval	26 Days	
DMP Development and Approval	12 Days	
DVM Development and Approval	17 Days	
Release of EDC into Production	39 Days	
<b>Total</b>	<b>44 Days</b>	

\*Business Days