

Is your Biotech/Medical Device Company Pre-Clinical?

Axiom can help accelerate your clinical trial program in critical ways.

Core Services

Data Management
Data Analytics
Clinical Operations
Project Management
Pharmacovigilance
Biostats

Pre-Clinical/Clinical Management

Clinical Consulting
Protocol Development
Study Plans Development
Regulatory Activities
SOP Development
Study Monitoring
Site Contracting/Budgeting
Risk Management
Clinical Monitoring
Medical Monitoring
Safety Monitoring
Vendor Selection/Management

Talk to our team of scientific & strategic specialists to see how we can help with your program plans, design, outcomes planning and so much more.

Getting a clinical program moving towards successfully planned and executed studies requires a holistic and well thought out approach. Connect with Axiom to find out how we can help make this a reality.

We help life science companies all over the world.

Chat with us to see how we assist in delivering on your eClinical and Study Operations needs.

650 Studies Completed. 18 Years.
Studies in over 30 Countries.



EDC | DM | IWRS | CTMS | ePRO

Data Analytics, eClinical Solutions & Services

Basics/Data Analytics

EDC/DM
Pre-Screening Log
Data Coding
General Log
Risk Management

ePRO/eCOA

ePRO/eDiary App
ePRO/eDiary Web Access
ePRO/eDiary Phone Access

Safety

AE/SAE Tracking
Safety Management

IWRS/RTSM

IWRS/IVRS
CTM Tracking

Data Import / Adjudication

Central/Local Lab Import
Imaging
Adjudication
Data Import
SAS On-Demand

CTMS

CTMS Dashboards
Deviations Management
Study Start-Up
Payment Tracking
Trip Reporting
eTMF



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axiom
real-time metrics™