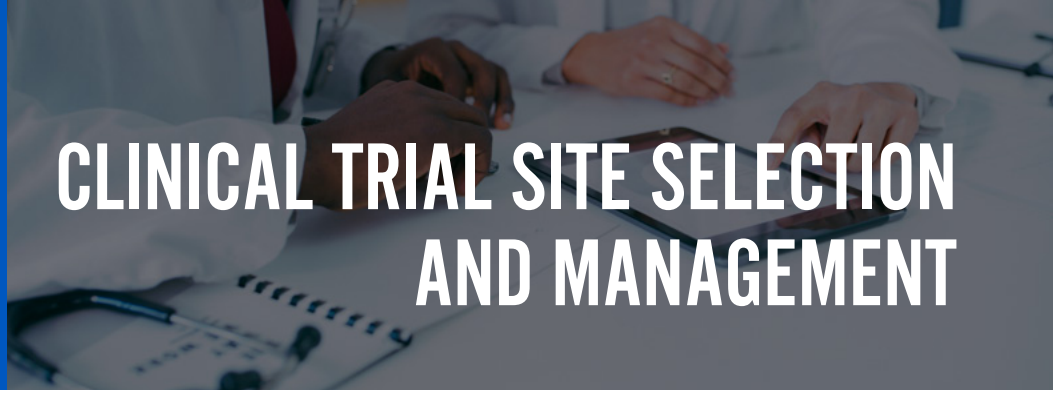
ALTASCIENCES



CLINICAL TRIAL SITE SELECTION AND MANAGEMENT

With more than 25 years of experience in early-phase drug development, Altasciences has conducted around 4,500 clinical trials in a variety of therapeutic areas at our three clinical pharmacology units. We leverage extensive data and experience to place your study where it makes the most sense, either at one of our clinical sites or at a partner site.

TAILORED CLINICAL TRIAL SITE IDENTIFICATION AND SELECTION

Collaborative Study Design
Our team will help you navigate the requirements needed to run your clinical trials smoothly and efficiently. Your Altasciences-appointed project manager will work directly with you, creating an open dialogue to align expectations against feasibility, identify prospective participants prior to site selection, and guide your study to completion.

Expert Site Identification and Feasibility Assessment
We thoroughly assess the capabilities of potential clinical sites based on your therapeutic area, geographic preference, and specialty population requirements. During the proposal stage, our physicians provide valuable protocol design feedback to address potential challenges.

Rigorous Vendor Assessment
Applying our internal standards and quality evaluations, we assess the experience, capabilities, therapeutic breadth, patient diversity, and risk factors of potential sites.

Comprehensive Analysis and Streamlined Site Selection
We fast-track your clinical trials by evaluating feasibility results and providing recommendations based on the number of required clinical sites and their locations. We have access to patient histories and data for quick recruitment, and take into consideration factors such as regulatory barriers, indication prevalence, and participant availability to adapt protocols for rapid and efficient study start-up.

Budget and Contract Negotiation
We will prepare for you a site budget comparison and ensure smooth clinical trial agreement negotiations—pre-established master clinical trial agreements with our top-tier strategic site partners guarantee expedited contracting.

CLINICAL SITE OPTIONS

With Altasciences, you have the option to utilize one or more of our three clinical sites, one of our affiliate sites, or both, for a **hybrid approach**.

Altasciences' Sites

Affiliate Sites

Hybrid Approach

contact@altasciences.com
altasciences.com

CLINICAL TRIAL SITE SUPPORT AND MANAGEMENT

Our highly experienced team of project managers ensures on-time and on-budget completion of your clinical trial projects, from Phase Ib to IIa. Every aspect is overseen as we meticulously monitor your studies' endpoints.

Extensive Partner Network
With a global network of over 100 specialized investigational sites, we have access to a large pool of study participants that enable you to take on even the most complex research projects. We leverage our connections to identify and qualify sites, build contacts, and manage your trial's performance, from start to finish.

Project Planning and Site Initiation
With a proactive, standardized, and integrated approach to planning, your dedicated cross-functional project manager will seamlessly guide you from one milestone to the next—to ease administrative burden and keep your studies on track.

Precision-Driven Management
We employ a distinctive partnering philosophy to ensure effective end-to-end collaboration, training, and resource planning for unparalleled support throughout your drug development process.

Dedicated project managers expertly oversee vendor selection and management, formulate comprehensive project plans and timelines, and facilitate seamless initiation of sites through coordination of start-up activities.

- **Efficient Investigator Grant Payments:** For financial peace of mind, we accurately manage your investigator grant payments in a timely fashion.
- **Effective Site Management:** We ensure adherence to protocols, on-time enrollment, and delivery of high-quality data.
- **Vigilant Trial Management:** We supervise study conduct and clinical monitoring, as well as lead support teams across the entire study timeline, supervising tasks from protocol development to preparing clinical study reports, whether managed in-house or through partner vendors.
- **Active Timeline Management:** We formulate comprehensive project plans and monitor timeline progression to ensure milestones are met.
- **Centralized and Proactive Communication:** Functioning as an extension of your own team, our project managers act as your primary point of contact to simplify your clinical operations.

contact@altasciences.com
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