Maximizing the Potential of Site-Centric Services

When to introduce Ergomed's Site Manager to your sites?

Site Management is Ergomed's optional service, introduced as an additional role to support investigational sites when it's most beneficial, such as:

- In larger, logistically more challenging clinical trials (multiple procedures, vendors, high volume of clinical visits).
- Particularly complex clinical trials where study procedures are outside of standard of care.
- Rare disease studies where site motivation & awareness needs to be maintained, patient recruitment & patient retention require additional initiatives.

We tailor our services!

Discover Ergomed's Distinctive Site Manager Role

Watch our recent webinar

"Maximizing the Potential of Site-Centric Services: Discover Ergomed's Distinctive Site Manager Role"

Supporting sites beyond the traditional monitoring model was the founding idea behind Ergomed.

Our site management support services are about keeping patients engaged and every data point is in check, predicting uncertainties and coping with the unknowns, alleviating any difficulties and delivering results. We take responsibility for the smooth conduct of your trial at the site.

Site Managers are an integral part of the study team and work closely with project management and CRAs to support the site staff (SNs, SSCs in particular). **Ergomed Site Manager provides mostly remote and less on-site support:**

- Support sites in first patient visit preparation
- Support in logistics of samples processing
- Assess patient compliance with visit attendance/treatment compliance
- Support in supplies ordering & tracking
- Facilitation shipments of samples
- Support in monitoring eDiary entries & compliance
- Assess site compliance in visit scheduling

Oncology Case Study:
Phase III, Head and Neck Cancer

Patient Recruitment Support

- Site-specific patient enrolment plans/goals,
- Feasibility of patient referral programs,
- Relationships with patient advocacy groups/communities,
- Implementation of agreed patient engagement procedures,
- Local/central advertising.

571 patients have been included through the Ergomed Site Management Model

Facilitating building confidence with site team leading to better control of study key risk indicators.

Ergomed Clinical Set-up:
Split of Responsibilities

Clinical Research Associate
Focus on Study Data and GCP Compliance

Standard Activities per GCP
Site Visits - Remote or Onsite
Remote capabilities - EMR where applicable, Regulatory records, IMP supply, lab information
Communication – Regular site contacts, deliverables, data quality
Risk & Issue Management – Early risk identification, management and follow up of issues
Data cleaning – Query Issuing, protocol deviation review
Safety Reporting/Consent Review

Site Manager
Logistical Support and Site Motivation

Logistic and Administrative Support - Patient Visit preparation, ensure study needs are met, supplies tracking, storage of supplies, sample processing, shipment of samples
Patient Recruitment/Engagement - Working with SC for Identification of patients, reduce SF rate
Patient Retention – Coordination of patient visits to reduce patient burden and increase patient retention
Continuous Site Training – SC/SN support & oversight (Lab processes, IMP handling, equipment); facilitating compliance

To learn more about how Ergomed's Site Manager Role maximize the value of site-centric services in your clinical trials, contact us at: info@ergomedplc.com

ERGOMED Clinical Research

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