IQVIA Study Coordinators and Study Nurse support services

Resources to support your clinical trial

Challenge

ADMINISTRATIVE WORKLOAD ON CLINICAL TRIALS TAKING TIME FOR PATIENT RECRUITMENT

Once your site is activated and ready to take part in a clinical trial, there's a lot of work that goes into setting up all study documents and getting organized for enrolling patients. In addition, finding and screening patients, can absorb also quite some time – work that's ideally delegated to a Study Coordinator. If you don't have someone designated and trained to handle these steps, your progress in recruiting patients is likely to suffer as you focus on patient care.

CONVENIENT, CAPABLE HELP WHEN YOU NEED IT

- Hold nursing degrees (not merely masters in biomedical sciences)
- Expert in all global Electronic Data Management systems
- Training in GCP
- Support Phase I-IV trials
- Skilled across therapeutic areas, including oncology and neurology
- Flexible arrangement to meet your changing needs

Solution

TRAINED STUDY COORDINATORS AT YOUR DEPARTMENT

IQVIA maintains a pool of study coordinators and study nurses who can be assigned to work at your site as contractors who are qualified to:

- Set-up subject visit worksheets
- Search your database to identify potentially eligible patients
- Complete Case Report Forms (CRFs)
- Perform study assessments (including vital sign assessments and blood draws

The terms of the placement are flexible to accommodate your changing resourcing needs. You could, for instance, book one coordinator for the duration of a trial or for the busiest recruitment months. Or you could arrange for help just certain days of the week or month. When the workload changes, so can your arrangement.

Results

The support that you get is tailored to your needs and invoicing is based on timesheets.

That allows you to **focus on patient care.**

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