SQC Clinical Trial
We care about your research

Keywords or phrase

Call us today on +44 (0)7913660781 and make sure your next clinical trial is run efficiently, professionally and safely.

SAFETY of your patients is our main concen. Avoid the cost of putting your patients health at risk and then delay the development of your new drugs. Place your study in the hands of a team with the strong commitment of protecting the patients safety & well-being.

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Ethics & GCP Forum, 13/Feb/2012 in London

SQC Clinical Trial representative will be present to the Ethics & GCP Forum organised by The Institute o...

Why an Increase of Regulatory Inspections in Clinical Trials?

The pharmaceutical industry is no doubt one of the most inspected. The reasons are obvious. Misconduct in re...

SQC Clinical Trial – We care about your research

Study and site management are two of the vital ingredients needed for your successful research trial.

By delivering a quality study within your timeline and budget, our professional knowledge and expertise combine to provide you with a service that meets your needs and expectations.

SQC Clinical Trial provides you with an innovative service with 3 main areas of focus:

Safety

The safety of your patients is protected at all times. Should any situation arise that could compromise the safety of your patients, we will inform you and the regulatory authorities immediately.

Quality

When developing medicines for your marketplace, it's essential your trial data is accurate. With SQC, the quality of your clinical data is assured. It will be accurate, complete, verifiable from source documentation and ready for inspection at any stage during the trial.

Communication

Effective communication from the outset is essential to ensure the smooth running of your clinical trial. By maintaining strong communication channels, from initial site investigations right through to the trial's conclusion, your patients will be recruited within timelines, your targets achieved and any protocol deviations avoided. This open approach will ensure there are no surprises along the way.

The SQC Approach

By concentrating on **S**afety, **Q**uality and **C**ommunication, we aim to eliminate delays in the development and availability of your drugs because ***we care about your research***.

Call us today and make sure your next study takes the SQC approach.

Registered Address: Freestyle House, 8 Mercia Business Village, Coventry, West Midlands, CV4 8HX, United Kingdom.

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SQC Clinical Trial – making your research our priority

Our Vision

The efficient setting up and running of your clinical trials is essential if they are to yield the quality of data you need.

Founded by Mylène Samuel, a freelance research scientist, SQC will offer you experience and excellence in the study and site management of your clinical trial.

Through continuous investment in training, SQC offers you:

- Efficiency of work delivered within your timeline and budget
- The ability to respond to changing markets
- The flexibility to provide you with a bespoke service
- Full safety of your patients
- Accurate and timely clinical data
- Effective communication
- Bi-lingual communication in English and French
- Full confidentiality

SQC Clinical Trial is based in the Midlands in the UK and has experience of working with sites in the UK, Ireland, France, Belgium and Switzerland.

Call us today and enjoy approachable and innovative support for your next clinical trial.

Our values

Adhering to strict guidelines, our aim is to provide you with exceptional service to exceed your requirements and expectations.

By focusing and adhering to Good Clinical Practice (GCP), EU directives, current national laws, regulations and your study protocols, we'll help you deliver market-leading medicines to patients around the world.

We understand GCP is essential within the clinical trials field as non-compliance results in poor data quality, a risk to the subjects' safety and the potential for your study data or marketing application to be rejected.

Mylène Samuel, representing SQC Clinical Trial, is a member of:
- The Institute of Clinical Research (ICR), the largest professional clinical research body in Europe and India
- Project Management Institute (PMI), therefore adheres to the Code of Ethics and Professional Conduct and the core values of responsibility, respect, fairness and honesty.

The Institute of Clinical Research

PMI Project Management Institute

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Clinical Trial Services

To maintain your reputation, it is essential your clinical trials are conducted accurately, within your timelines and on budget.

Our fresh, modern approach and range of clinical trial services ensure you receive a professional service for each of your trials. Plus, our professional knowledge and expertise ensure we fully understand your requirements so we always exceed your expectations.

Below, you will find the 3 main areas of our clinical trial services. If you need any further information, or would like to talk further, please get in touch and we'll be glad to help.

Study start-up management

• Developing and writing Clinical Trial protocols to outline the purpose and methodology of your Clinical Trial

• Presenting Clinical Trial protocols during meeting

• Designing data collection forms (Case Record Forms)

• Co-ordinating with the Ethics Committee, which safeguards the rights, safety and well-being of all Clinical Trial subjects

• Managing regulatory authority applications and approvals (that oversee the research and marketing of new and existing drugs)

Site management

• Locating and assessing the suitability of facilities at a study centre

• Briefing doctors/consultants (or investigators) on conducting Clinical Trials

• Setting up the study centres – ensuring each centre has the Clinical Trial materials, and training site staff to Clinical Trial specific and industry standards

• Monitoring the Clinical Trial throughout its duration, which includes visiting the study centres on a regular basis

• Auditing clinical trial procedures, including regulatory affairs and good clinical practice

• Verifying that data entered on the Case Record Forms is consistent with patient clinical notes (Source Data/Document Verification)

• Collecting completed Case Record Forms from hospitals and General Practices

• Writing visit reports

• Filing and collating Clinical Trial documentation and reports

• Ensuring all unused Clinical Trial supplies are accounted for

• Closing down study centres on completion of the Clinical Trial

• Archiving study documentation and correspondence

Study management

• Conduct accompanied visits and review visit reports

• Study team training and support

• Study data tracking

• Review of the study's progress

• Study budget and timeless tracking

• Archiving study documentation and correspondence

• Preparing final reports

By working with SQC Clinical Trial, you'll be working with a team with a wide range of industry experience, including the areas of Oncology, Cardiology, Haematology, Infectious diseases and Central Nervous System.

Call us today and make sure your next clinical trial is run efficiently, professionally and safely.

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