

Industry Visit: Write-up

The Site Visit serves as an invaluable opportunity to gain comprehensive insights into the dynamic front-end activities of a clinical trial environment. It offers a unique vantage point for Investigators, Researchers (including clinical research coordinators), Ethics Committee members, and Clinical Trial Participants to engage in a holistic exploration.

During this enlightening visit, attendees will have the privilege to:

- I. Engage in meaningful dialogues with investigators, site personnel fostering a deep understanding of the pragmatic challenges they encounter. These challenges encompass identifying suitable patients, persuading them to partake in clinical trials, administering investigational medicinal products (IMPs), conducting localized laboratory and diagnostic tests, evaluating study outcomes, and adeptly utilizing electronic Case Report Forms (eCRFs) and provided technologies.
- II. Immerse themselves in the sphere of the research team stationed at the site. This team adeptly handles non-medical facets of clinical trials, capturing source data, entering information into eCRFs, surmounting real-world hurdles in addressing eCRF queries, and skillfully fostering engagement and retention among Clinical Trial Participants.
- III. Engage with the esteemed members of the Ethics Committee, entrusted with the responsibility of meticulously scrutinizing study documents from both scientific and ethical lenses. Their pivotal role involves granting approvals, as well as astutely evaluating the causal relationships between Serious Adverse Events (SAEs) and participation in the clinical trial.
- IV. Interface with the integral figures in the clinical trial landscape — the Clinical Trial Participants themselves. This interaction unveils the intricate balance between potential benefits and inherent risks that their participation entails, thereby enabling them to make informed decisions about their health and well-being.

In addition to the above learning, you may also foster connections with the visionary leadership of the Clinical Trial Site, including Dr. Ferzaan Engineer, Co-Founder & Chairman, Suresh Ramu, Co-Founder & CEO and Amit Sharma, CIO & Head PartnershipS & CSR, contingent upon their availability. This rare opportunity offers a glimpse into the minds shaping the site's strategic direction and underscores their dedication to advancing the realms of clinical research.

In summary, the Site Visit transcends a mere observational experience, transforming into a multidimensional journey of enlightenment, engagement, and empowerment. This visit offers an exclusive opportunity for the participants to enhance their perspectives, amplify understanding, and invigorate collaboration across the diverse spectrum of clinical trial stakeholders.

Registration cost: INR 1000+Taxes