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Senior Specialist - Clinical and Ancillary Supplies Services

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Company: Fortrea

Location: Buenos Aires

Category: other-general

The Clinical & Ancillary Supplies Services (CASS) Senior Specialist:

Coordinate and oversee aspects of logistics related to Clinical and Ancillary Supplies services.

Support these services to both internal and external clients.

Responsible for the set-up and coordination of project activities related to clinical and ancillary supplies

Independently support the delivery of the Clinical and Ancillary supplies component of assigned studies within a region or for a large global study.

Provide local support with clinical and ancillary supplies start up activities and related project feasibility

Act as a subject matter expert (SME) related to import, export and labeling requirements.

Work with staff in Clinical Trial Design & Strategy, Vendor Management, Quality Assurance, Business Development and Operations, to provide relevant clinical and ancillary supply data critical to the ability of the organization to develop plans for the successful implementation and conduct of global clinical trials.

Assure that pre-study, drug supply and labeling processes comply with applicable guidelines and regulations and fulfill client and Covance requirements.

Provide logistical support for clinical and ancillary supply coordination

Provide input into business development activities

Act as a subject matter expert (SME) in client focused clinical and ancillary supply workstreamsy

Required:

Minimum of 4 years relevant clinical research experience in a pharmaceutical company/CRO, including at least 2 years of clinical and ancillary supplies management responsibility.

University/College Degree (preferably postgraduate degree)

Fluent English

Very strong technical writing skills and analytical skills

Strong networking abilities and an ability/willingness to work with individuals across the globe

Excellent communication, planning & organizational skills

Ability to inspire, to lead by example and motivate teams to seek solutions and get results

Strong abilities to negotiate and liaise with customers in a professional manner

Experience in end to end (full service) clinical trial management, ideally with global project management experience

Abilities to review/analyze relevant data so as to build evidence-based clinical trial strategies

Abilities to design/develop effective strategies to bring about trial success, as defined by clients and project teams through the use of an evidence-based approach.

Strong computer skills with an ability to understand and access and leverage technology alternatives.

Ability to work independently

Availability to travel (10-20%)

Preferred:

2-4 years Clinical Supply Project Management logistics experience

Demonstrated Regulatory, GMP or QA experience.

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