Veronica Cisneros
Curriculum Vitae

NAME:
Veronica Cisneros

October 2021 - Present

ICRA - PAREXEL SA

The Clinical Research Associate is the clinical sites' direct point of contact and accountable for managing site quality and delivery from site identification through close-out. The aim of this role is to build the relationship with the site, to be the sole position accountable for the site performance, including driving the site successfully to initiation, the development of a robust patient recruitment strategy, setting up a system, assessing compliance with regulations and the site's capability to comply with the study needs and recruitment expectations, and problem solving to address and resolve site issues.
Act as company direct point of contact with assigned sites, accountable for quality and delivery during the start-up phase.

June 2018 - October 2021

Clinical Research staff at Centro de Estudios Infectológicos Stamboulian Health Services
Clinical Research staff at Infectious Diseases Division in Fleming Institute

Clinical Research Experience
Sub investigator:

June 2014 - March 2020. Clinical trial Division Stamboulian Health Services

Performing study–related procedures and /or to make important study-related decisions in compliance with the ethical conduct of the study, submitting new protocols, protocol amendments, continuing reviews, and safety reports to the appropriate IRB for review, maintaining regulatory binders in accordance with sponsor specifications and general industry standards under the supervision of the PI.

- supporting the Principal Investigator during Bioresearch monitoring (BIMO) inspection conducted by the US Food and Drug Administration.
- supporting the Principal Investigator during ANMAT (Administración Nacional de Medicamentos, Alimentos y Tecnología Médica) audit.

Study Coordinator:

March 2020 - October 2021. Clinical trial Division Stamboulian Health Services

Performing activities such as contact with sponsors/CROs, Central Labs, IEC (preparation and submission of documents). Completing CRFs, organizing logistics for patient's visits, participating in site initiation visits. Contract negotiations.

Study Manager

Jan 2021 - Oct 2021. Pharmexx Arg

Performing activities and providing accountability for day-to-day operations of the project, as defined by the contract and according to ICH/GCP and all other applicable laws, rules, and regulations, reviewing and provide input for study protocol, edit check specifications, data analysis plan, and final study report, developing operational project plans, customizing documentation for sites to submit to regulatory authorities.

EDUCATION:

University Education

Universidad Nacional del Nordeste. School of Medicine. Buenos Aires. Argentina.
Degree: Medical Doctor

Postgraduate Training

Internal Medicine Residency. Hospital de Clínicas “José de San Martín”.
Degree: Internal Medicine Specialist.

Chief Resident. Internal Medicine Residency. Hospital de Clínicas “José de San Martín”.

Infectious Disease Fellowship. FUNCEI (Fundación del Centro de Estudios Infectológicos).

Course Infectious Diseases in Adults 2015. Harvard Medical School. Boston – EUA.

Infectious Disease Specialization. CEMIC (Centro de Educación Médica e Investigaciones Clínicas)
Degree: Infectious Disease Specialist

Clinical Research Fellowship. FUNCEI (Fundación del Centro de Estudios Infectológicos).

Master's Degree in Clinical Effectiveness. Instituto de Efectividad Clínica y Sanitaria (IECS)
Magister Candidate (MSc (c)). Ciudad Autónoma de Buenos Aires (CABA)

Published Scientific Papers

Sex Transm. Infect. 2016 Nov 17. pii: sextrans-2016-052893. doi: 10.1136/sextrans-2016-052893. Very high incidence of syphilis in HIV-infected men who have sex with men in Buenos Aires city: a retrospective cohort study. Bissio E, Cisneros V, Lopardo GD, Cassetti LI. https://www.ncbi.nlm.nih.gov/pubmed/27856515

LANGUAGE SKILLS:

English: Fluent - Able to perform translations and validations
Spanish: mother tonge