Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

EDUCATION

2021 (ongoing)	Medical Affairs in Medicines Development Professional Certification Program ProgramDescriptionIFAPP Academy_King's College London
2006	Postgraduate Diploma in Clinical Pharmacological Investigation Instituto de Biología y Medicina Experimental (IBYME), Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET), Universidad de Buenos Aires (UBA), Argentina
2001	Physical Therapist Universidad del Salvador CABA, Argentina
1996	Bachelor Colegio Nuestra Señora de Huerto, Pergamino, Buenos Aires, Argentina

PROFESSIONAL EXPERIENCE:

April 2019 to present: Medical Science Liaison
Ophthalmology Franchise
Novartis Argentina S.A

- Development peer to peer scientific relationships, assess advocacy (degree of Medical Expert understanding and agreement with medical platform statements) and contributions (activities in which Medical Experts are independently qualified to engage or collaborate with the organization), and respond to unsolicited requests for scientific information using approved, non-promotional materials with Medical Experts/Key Stakeholders, Health Care Professionals (HCPs) and decision makers.
- Contribute to identification and mapping of Medical Experts/key stakeholders in line with local strategies.
- Provide and discuss scientific information and data with HCPs to ensure quality and accuracy of medical and scientific information on new treatment options and selected areas of therapeutic interest.
- Liaise and provide up-to-date medical support to relevant
- External groups to ensure evidence based understanding of relative to medical guidelines, and health economic information.
- Provide speaker training to HCPs to support education to the healthcare community about new therapies developed. cientific Research Support
- Provide scientific educational activities to support study accrual for key trials as appropriate.
- Provide feasibility and research site recommendation, sponsored clinical trials.
- Collaborate with R&D team and Medical Advisors, to support clinical sites, including providing guidance in accordance with guidelines,, and regulations as appropriate.

Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

- Increase awareness clinical research program to enhance patient identification and trial accrual
- Upon investigator request, liaise between potential researcher and the Company on IIT submission process or requests for research grade substance (Material Transfer Agreements) to support independent clinical/preclinical research. Scientific Exchange/Insights
- Engage in scientific discussions related to research areas/compounds/pipeline with MEs, speakers & investigators.
- Support scientific exchange to advance understanding of new scientific principles, novel research trends, and current scientific debate.
- Provide scientific and educational information to healthcare professionals on specific patient critical issues.

January 2017 to March 2019: Expert Clinical Research Associate-CRA III:
Novartis Argentina S.A

Central point of communication/contact between Novartis and the Investigator site personnel to ensure:

- That the trial relates activities are conducted at site following GCP and Novartis standards (processes and SOPs)
- That agreed targets (e.g. number of patients, key milestones) are met.

The expert CRA is able to work without supervision on complex Trials. Facilitates and support the local/cluster study team to achieve strategic objectives through the efficient execution of allocated trials.

December 13 to December 2016: Sr. Clinical Research Associate
Inventiv Health Clinical

- Perform routine site visits, including pre-study, initiation, interim, and closeout visits. Visit to include monitoring of proper informed consent procedures, compliance with protocol, GCP/ICH Guidelines and other applicable regulatory requirements and assurance of good site performance. This is accomplished by detailed review of subject records, essential documents, investigational product disposition and accountability, site personnel and procedures.
- Manage assigned sites by regular contacts to ensure site compliance, adequate enrollment, and understanding of study requirements.
- This is accomplished by completion of monitoring reports and follow-up letters within the project-specific timelines. Significant issues noted must be reported to appropriate personnel immediately.
- Maintain project tracking system of subject and site information.
- Assist with company's quality control initiative.

Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

- Support project management team with assessment of workload and site assignments within the project team.
- Participate in company-required training programs.

Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

Participate in a Trial Master File Audits and Sponsor Audits.

- Perform necessary administrative functions (e.g., tracking of expense reports, time and attendance).
- Maintain home office (e.g., procurement of office supplies, submission documents), if applicable.

March 11 to November 13

Clinical Research Associate II
i3 Pharma Resourcing

- Perform routine site visits, including pre-study, initiation, interim, and closeout visits. Perform responsibilities with minimal support from management. Visits to include monitoring of proper informed consent procedures, compliance with protocol, GCP/ICH Guidelines and other applicable regulatory requirements, and assurance of good site performance. This is accomplished by detailed review of subject records, essential documents, investigational product disposition and accountability, site personnel and procedures.
- Manage assigned sites by regular contacts to ensure site compliance, adequate enrollment, and understanding of study requirements.
- Report to i3 project team, client, and site personnel any findings noted at monitoring visits.
- This is accomplished by completion of monitoring reports and follow-up letters within the project- specific timelines. Significant issues noted must be reported to appropriate personnel immediately.
- Maintain eClinical or alternative project tracking system of subject and site information.
- Participate in a Trial Master File Audits and Sponsor Audits.
- Participate in company-required training programs.
- Perform necessary administrative functions (e.g., tracking of expense reports, time and attendance).
- Maintain home office (e.g., procurement of office supplies, submission of documents), if applicable.

April 2008 to March 2011

Clinical Research Associate I
Kendle International

- Performed routine site visits, including pre-study, initiation, interim, and closeout visits. Visits to include monitoring of proper informed consent procedures, compliance with protocol, GCP/ICH Guidelines, and other applicable regulatory requirements, and assurance of good site performance.
- Managed assigned sites by regular contacts to ensure site compliance, and understanding of study requirements.
- Reported to project team and site personnel any findings noted at monitoring visits. This was accomplished by completion of monitoring reports and follow-up letters within the project-specific timelines.
- Performed Start up activities.

Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

- Participate in a Trial Master File Audits and Sponsor Audits.
- Maintained project tracking system of subject and site information.
- Participated in company-required training programs, as well as, in Sponsor Investigator Meetings.
- Performed necessary administrative functions.

May 2007 to March 2008

Project Assistant
Kendle International.

- Provided Administrative support to the CRAs in diverse in house activities such as filling, Sites contact and study correspondence, kept updated Tracking of Lab results, patient enrollment and study queries.
- Performed Regulatory activities such as submissions to MoH and IRBs.
- Inventoried and maintenance of office materials and equipment.
- Participated in a study close out.
- Maintained project tracking system of subject and site information.
- Participated in company-required training programs
- Performed necessary administrative functions

Jan 2002 to May 2007

Physical Therapist
Kinesiology Institute
CABA, Buenos Aires, Argentina

- Physical therapist Staff Coordinator.
- Specialized in attention to Patients in Trauma and Respiratory area.

FOREIGN LANGUAGES

Spanish — Native language
English — Advance
New England School of English, Boston USA.
January 2008

SUMMARY OF PROJECT EXPERIENCE

Therapeutic Area, Sub Indication and Drug Class	***Phase***	***Job Title***
Inmunology/ Renal Transplant	II	Lead CRA/Expert CRA III
Inmunology/ Lupus Nephritis	II	Lead CRA/Expert CRA III
Rheumatologis/Ankylosing Spondylitis	IIIb	Lead CRA/Expert CRA III
Inmunology/ SSP	IIb	Lead CRA/Expert CRA III
Inmunology/ SLE	II	Expert CRA III
Hepatology/ NASH	II	Expert CRA III
Inmunology /Psoriatic	III	Expert CRA III
Inmunology/Hip Fracture	IIa/IIb	Expert CRA III
Immunology/Renal Transplant	IV	Expert CRA III
Immunology/Children Renal Transplant	IV	Expert CRA III

Valeria Ferreiro
Physical Therapist – USAL

Cel Phone: +5491131142087

Mail: valeferreiro79@gmail.com

Rheumatologist/Psoriatic arthritis	III	Expert CRA III
Respiratory/ Asthma	III	Sr. CRA
Respiratory/ Pulmonary hypertension	III	Sr. CRA
Respiratory/ Children Pulmonary hypertension	III	Sr. CRA
Respiratory/COPD	IV	CRAII
Cardiology/hypertension	III	CRA
Cardiology/ resistant hypertension	III	CRA
Rheumatologist/Lupus	II	CRA
Rheumatologist/rheumatoid arthritis	II	CRA
Rheumatologist/Lupus	III/observational	CRA
Regulatory/ audit file	Audit file	CRA
Nephrology/epidemiology study	IV	PA
Women health/ Protocol evaluation of the predictive ability of angiogenesis factors for preclampsia.	IV	PA