Lic. Luciana Mannarino
Lugar de residencia: Lanús Este.
Tel: 15-5420-9471
Email: **luciana.mannarino@hotmail.com**

Buenos Aires, enero del 2021

De mi mayor consideración:

Tengo el agrado de dirigirme a Ud. a fin de solicitarle tenga a bien considerar el curriculum adjunto ante una posible vacante en las búsquedas laborales que su empresa estime pertinentes.

Es de gran interés para mí trabajar y desarrollarme en empresas de vuestro renombrado prestigio. Me siento capacitada para desempeñarme en áreas que demanden un alto compromiso y flexibilidad, dado que he tenido la oportunidad de adquirir experiencia poniendo a prueba mi idoneidad y capacidad.

Quedando a vuestra entera disposición para concretar una entrevista personal en fecha y horario que Ud. estime conveniente, hago propicia la oportunidad para saludarlo cordialmente.

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Personal Details:

Date of birth: 2nd, March 1989.

Nationality: Argentinian.

ID: 34390514

Code Post: 1824

Address: O´ Higgins 1557- 1A-Lanus Este.

Mobile phone: 15-5420-9471

E-mail: luciana.mannarino@hotmail.com

CUIL: 27-34390514-4

License Nutritionist Number:

-National: 7215

-Provincial: 2700

Work Aim:

Develop myself optimally in the role assign to me and develop my professional skills.

Studies/ Qualifications

-1997/2006: English Study- Asociación de ex alumnos Del Profesorado en Lenguas Vivas "Juan R Fernández".

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-**2006:** Graduated from the Institute José Manuel Estrada de Lanús- Degree of Bachiller en Economía y gestión de las organizaciones.

-**2007**: CBC (Biochemistry orientation) in Buenos Aires University.

-**2008**: First quarter at College of Pharmacy and Biochemistry in Buenos Aires University.

-**March 2008**: Approval of the international Exam "First Certificate in English".

-**2009-2012**: Graduated from the career Nutrition in Maimonides College, obtaining a Diploma of Honor for having the best career average.

Language:

-English: speaking, reading and written (advanced)
-Portuguese: speaking, reading and written (basic).

Technical knowledge:

Pc- Pack office.

Work experience:

Date of Employment: 02/2012- 10/2015
Job Title: Food security.
Name of Employer: YPF
Key Responsibilities:
-Making audits about Quality and GMP (Good Manufacturing Practice).
-Taking part in the creation of the checklist that were used to supervise the Convenience Store.
-Giving training about GMP to the employees.
-Performing reports with the results of the quality audits and sending it to different areas of the company.
-Sending the protocols with the results of food and water analysis.
-Meeting food suppliers and visiting their factory in order to make a check-up.

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Date of Employment: 10/2015 - 03/2016
Job Title: CRA Trainee- Clinical Research Associate.
Name of Employer: QUINTILES
Key Responsibilities:
-Complete appropriate therapeutic, protocol and clinical research training to perform job duties.
-Gain experience in study procedures by working with experienced clinical staff in Clinical Research Associate (CRA), Clinical Research Specialist (CRS) and Clinical Trials Assist (CTA) roles.
-Collaborate with other functional groups such as Data Management and Pharmacovigilance.

Date of Employment: 04/2016 - 07/2017
Job Title: CRA 1- Clinical Research Associate.
Name of Employer: Quintiles IMS (Ex Quintiles)
Key Responsibilities:
-Perform site selection, initiation, monitoring and close-out visits in accordance with contracted scope of work and good clinical practice.
-Support development of project subject recruitment plan on a per site basis.
-Work with sites to adapt, drive and track subject recruitment plan in line with project needs to enhance predictability.
-Provide monitoring visits and site management for a variety of protocols, sites and therapeutic areas.

Date of Employment: 08/2017 – 05/2018
Job Title: CRA 2- Clinical Research Associate.
Name of Employer: IQVIA (Ex: Quintiles)
Key Responsibilities:
-Administer protocol and related study training to assigned sites and establish regular lines of communication with sites to manage ongoing project expectations and issues.
-Evaluate the quality and integrity of study site practices related to the proper conduct of the protocol and adherence to applicable regulations.
-Escalate quality issues as appropriate.
-Manage the progress of assigned studies by tracking regulatory submissions and approvals, recruitment and enrollment. May support start-up phase.
-Create and maintain appropriate documentation regarding site management, monitoring visit findings and action plans by submitting regular visit reports, generating follow-up letters and other required study documentation.
-Build awareness of features and opportunities of study to site.
-Collaborate and liaise with study team members for project execution support.

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Date of Employment: 05/2018 – 09/2020
Job Title: CRA 2- Clinical Research Associate.
Name of Employer: Abbvie
Key Responsibilities:
-Administer protocol and related study training to assigned sites and establish regular lines of communication with sites to manage ongoing project expectations and issues.
-Evaluate the quality and integrity of study site practices related to the proper conduct of the protocol and adherence to applicable regulations.
-Escalate quality issues as appropriate.
-Manage the progress of assigned studies by tracking regulatory submissions and approvals, recruitment and enrollment. May support start-up phase.
-Create and maintain appropriate documentation regarding site management, monitoring visit findings and action plans by submitting regular visit reports, generating follow-up letters and other required study documentation.
-Build awareness of features and opportunities of study to site.
-Collaborate and liaise with study team members for project execution support.

Date of Employment: 09/2020- present
Job Title:
Senior CRA - Clinical Research Associate.
LSR: Local Study Representative

Name of Employer: Abbvie
Key Responsibilities:

-Serves as the country SMM primary point of contact accountable for a specific study.
-Ensure successful protocol level execution of SMM deliverables involving site selection, start-up, execution, and close-out.
-Proactively identifies and mitigates site issues that could affect study deliverables.
-Oversee site/central EC submission and approval timeline.
-Support contract negotiation process.
-Oversee IP importation process.
-Oversee any local specific approval process.
-Review systems for logic and completeness (i.e. site assignments, event dates, submissions/approvals) and follow up with CRAs for any updates or corrections
    - Communicate early and often to CRAs about expectations for entry of required fields
    - Follow-up on emails highlighting missing data for a site
    - Identify trends and/or delays across country sites (i.e. Extended CSA execution and/or sites not receiving needed information)
-Liaise with CRAs about non-responsive sites and/or delays resulting from site activities for appropriate follow-up (i.e. budget/contract negotiation issues, missing and/or incomplete documents needed for IP Authorization)
-Review Monitoring Plan
-Provide guidance to SMM team to expedite processes/tasks that affect timely start-up for sites, including:

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- Communicating the country site activation plan with defined timelines
- Completion of study-specific training and/or approvals required for activation, such as submission of test images to vendors and completion of training for unblinded/blinded staff via portals.

-Serve as a training mentor for other LSRs. Serve as a role model, guide and mentor for newly-assigned LSRs and others LSRs (mentoring new-LSRs in other countries).

Seminars:

- Seminar "Probiotics". Dr. Tomas Herrero; Ing. Ricardo Weill; Dra. Eugenia Maciero; Lic. Gabriela Olagnero. Maimónides University (June 2010).
- Seminar "Food Research: Research Models Applied in Foods" Maimónides University. (October 2011).
- Working Day "Industry and Gluten-free Food. Treatment of Celiac disease". German Hospital. Area of Hygiene and Food Safety (5 hours) (May 2012).
- Working Day "Knowing our Health". Maimónides University (16 hours) (September 2012).
- Technical Training Program. "Basic Security Module". YPF. (September 2012).
- Advanced Training For Internal Trainers. YPF. Graciela Garone. Management consulting (16 hours) (October 2012).
- Food Handler. YPF. Government of Buenos Aires (December 2012).

Congresses and Conferences:

-XXXV Annual Meeting. Argentine Chapter of the Latin American Society of Nutrition (CASLAN). VIII working day of Nutrition and Food Technology. Lujan University. September 2012.

Investigation Work:

-"Survey levels of Arsenic, Nitrites and Nitrates in drinking water in different geographical areas of Argentina". Authors: Mannarino Luciana. Marchini Mariano. Qualification 10 (ten). September 2012.

-"Advertising messages of food and beverages in children`s TV channels". September 2012.

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Postgraduate Courses:

-**Pharmacology and Nutrition**. "Argentina Society of Obesity and Eating Disorders" – SAOTA. (182 Teaching hours).

-**Management Nutritional Consultation: Cardio-metabolic Clinical Cases**. "Nutrition Society Argentina"-SAN. (220 Teaching hours).

-**Good Manufacturing Practices (GMP) and Food Services Management**. "Argentina Association of Dietitians and Nutritionists"- AADYND. (184 Teaching hours).

Other Relevant Information

Licenses and Certifications

- GCP Barnett Accreditation, 2016
- Oncology Clinical Research Fundamentals, 2016
- Monitoring Fundamentals, 2016
- Methodology of Research and Monitoring of Studies, 2017

Certificates are available for all courses, conferences, jobs and postgraduate courses.

Electronically signed by: Luciana Mannarino
Reason: Approver
Date: Jan 24, 2021 18:49 GMT-3