

EVANGELINA BREDA

B.S of Molecular Biology in Medical Sciences

OUTSTANDING SKILLS

- Conflict resolution
- Proactive attitude
- strong work ethic
- Search for continuous excellence.
- Great attention to detail.
- Great ability to adapt to changes.
- Great capacity for clear and assertive interaction with the different stakeholders.
- Analytical capacity with Honesty and courage.
- Innovative and inclusive spirit
- growth mindset

CONTACT INFO

(+54) 1155152455

evangelinabreda@gmail.com

Buenos Aires, Argentina

EDUCATION

UNIVERSIDAD FAVALORO
2008 - 2013
B.S of Molecular Biology in Medical Sciences

UNIVERSIDAD AUSTRAL
2013 - 2014
Design, methodology and monitoring of clinical trials

INSTITUTO TECNOLOGICO BUENOS AIRES (ITBA)
2013 - 2014
Business management for scientists and technologists.

CODER HOUSE
Junio 2021 - Diciembre 2021
Product Manager

OTHER

Language
English as a second language (Fluent)
Italian as a foreign language (Conversational, intermediate)

Software
Office (Superior)
CRM (Intermediate)
Veeva Bolt (Intermediate)

WORK EXPERIENCE

BIOPHARMA QUALITY OPERATIONS SR. MANAGER (COMMERCIAL PHARMACOVIGILANCE MANAGER)

Pfizer Argentina | APR 2017 - present

Descripcion:

Collaboration with the Business department with the management and monitoring of the program Patient support programs, awareness activities, primary market research and business development.

Design and implementation of projects associated with interactions with external clients, where a two-way communication is opened , providing the added value of compliance with regulations, ensuring the commercial interests of the organization focused on the needs of each business unit.

Direct interaction with the main Stakeholders (medical, Regulatory, Legal, Marketing, Commercial, Compliance) of the local market in the follow-up of commercial decisions in relation to new challenges focused on innovation, well-being of patients and clients and their needs through their Patient journey and customer experience.

Active participation in E2E projects of Local, Regional and Global scope leading multidisciplinary teams (Regulatory, Medical Affairs, Legal, Commercial, IT, Digital strategists, Marketing and product development, etc).

Development of effective strategies focused on the needs of the business, valuing those of the external client (patients, health professionals, etc.), by identifying pain and opportunities for improvement in their experience (Patient Journey / Customer journey) from the moment of diagnosis to access to the appropriate treatment (according to business unit and pathology) and its follow-up.

Leadership in consultations and risk analysis (compliance), continuous improvement and good practices focused on product and patient safety (pharmacovigilance).

Interaction between multifunctional teams and coordination of activities that impact each of them and their KPIs. Creation of work plans and ownership of KPIs to define priorities, pain points through the coordination and incorporation of innovative strategies.

Preparation and conduct of internal and external audits (vendors) to ensure compliance with processes related to the management of information based product and patient safety (Quality Assurance).

Implementation and monitoring of processes focused on local and global regulations and internal policies. Quality assurance by design of quality checks, audits and effectiveness checks.

Audit and inspection readiness activities implementation and follow up. CAPA plans design, implementation and follow up.

LOCAL PHARMACOVIGILANCE MANAGER - ARGENTINA, URUGUAY & PARAGUAY.

Boehringer Ingelheim Argentina | Nov 2014 - Mar 2017

Local responsible for pharmacovigilance in front of the local regulatory authority (ANMAT).

Management of detection, reporting and analysis of spontaneous and requested adverse events.

Development of documentation for submission of new products and update of security profile (pre and post commercialization).

Design and implementation of clinical trial monitoring plan.

Monitoring of commercial initiatives associated with the interaction with Health Professionals, Patients and consumers.

Quality assurance and compliance with internal processes (internal audit).

Quality assurance and compliance with local and global regulations (Inspection readiness).

Risk analysis focused on product safety.

Internal and external Audit & inspection readiness participation. CAPA plans design, implementation, monitoring and effectiveness checking

CLINICAL RESEARCH PROJECT COORDINATOR

IQVIA (ex – QUINTILES), Argentina | Jan 2014 - Dec2014