



Curriculum Vitae

Personal information **Janice Cassar Giacomotto**

Work experience

1. Employer: Medicines Authority
 - Start date: 012018
 - End date:
 - Position: Senior Medicines Inspector
 - Activities: Mainly responsible for carrying out GMP, GDP and Pharmacovigilance inspections. I also represent the Medicines Authority in WGEO meetings
 - Country: Malta
2. Employer: Combino Pharm Malta
 - Start date: 112017
 - End date: 012018
 - Position: Quality Management Officer
 - Activities: Routine QA activities
 - Country: Malta
3. Employer: Starpharma Ltd.
 - Start date: 122003
 - End date: 112017
 - Position: QA Manager and QP
 - Activities: Apart from the day to day running of the QA Department, my responsibilities included: • Ensuring that the quality system of the company was according to GMP, • Approving SOPs, QC documentation, batch records and quality documents • Overseeing the approval of suppliers. • Writing and/or approving PQRs • Co_ordinating risk assessments related to quality issues. • Carrying out internal and external audits as well as compiling related reports and CAPA plans • Providing Induction and GMP training to new employees • Assisting clients with the registration of products and variations. . I was also responsible for replacing the main QP in his absence.
 - Country: Malta
4. Employer: Malta National Laboratory
 - Start date: 082002
 - End date: 122003
 - Position: Forensic Scientist
 - Activities: I was employed as a forensic scientist, where I was trained as a fingerprint expert. I also used to help in the chemistry division where I gained practical laboratory experience. I was responsible for analysing different materials such as textiles, food and seawater. Furthermore, I worked on a 5th Framework European project (NITECRIME), which involved the use of ICP/MS for the analysis of trace elements.
 - Country: Malta

Education and training

1. Subject: University of Manchester
 - Start date: 062009
 - End date: 062012
 - Qualification: MSc. Industrial Pharmaceutical Sciences
 - Organisation: Modules studied include Pre_formulation, Quality Control and Quality Assurance, Regulatory Affairs, Laboratory Testing, Solid Oral Dosage Forms, Pharmaceutical Engineering and Product Development. Thesis: Training aspects in the manufacture of solid oral dosage forms
 - Country: United Kingdom
2. Subject: University of Malta
 - Start date: 101997
 - End date: 062002
 - Qualification: B. Pharm Hons
 - Organisation:
 - Country: Malta

Additional information

Publications

Projects

Memberships

Member of the Malta College of Pharmacy Practice, Member of the Malta Pharmaceutical Association, Member of the Malta Qualified Persons Association, Member of the European Qualified Persons Association

Other Relevant Information