



José Malta, PharmD, MSc

Executive Director and Head of Regulatory Affairs at Nutri.add – Healthcare

Advanced Solutions, S.A.

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Curriculum Vitae Synopsis

Summary

With a background in Pharmaceutical Sciences and a continuous training aiming the updating and deepening of required subjects, works in the field of scientific and regulatory affairs of medicines, healthcare and nutritional products for the last 10 years.

Experience

Product Development Coordinator at Nutri.add S.A. | Group Labialfarma (*nutraceuticals, herbal products, medicinal products, medical devices, veterinary products, cosmetics and biocides*). | Pharmaceutical consultant, Regulatory Affairs, Quality Assurance, Pharmacovigilance project manager and department's coordinator, HR responsible, customer service and business development at Phagecon – Pharmaceutical Consulting and Services, Lda. | Technical Responsible of Orthopaedics Division at Smith&Nephew Lda. | Regulatory Affairs officer at Mepha, Lda. | Trainee in MKT, Sales, Regulatory Affairs and GMP at GlaxoSmithKline Consumer Healthcare Ltd. | Trainee in Scientific and Regulatory Affairs at Phagecon, Lda. | Trainee at Community and Hospital Pharmacies. Trainee at Laboratory of Pharmacognosy at Faculty of Pharmacy of University Louis Pasteur (Strasbourg).

Education

Graduated in Pharmaceutical Sciences from University of Coimbra, Faculty of Pharmacy. MSc at Faculty of Pharmacy of University of Coimbra. Post-Graduation course in Law of Pharmacy and Medicines from University of Coimbra, Faculty of Law. MSc Student of Pharmaceutical Medicine at University of Aveiro. Title of Specialist in Regulatory Affairs granted by Portuguese National Order of Pharmacists.

Some additional related training to be highlighted: EudraVigilance – electronic reporting of ICSRs | DIA/EMA/INFARMED IP; Jornada Extractos Vegetales: Extractos Vegetales en la Industria Cosmética y Alimentaria: Normativa, Calidad y Avances | Universidad Alcalá - Madrid; Technical meeting on the reporting of human studies submitted for the scientific substantiation on health claims | EFSA Nutrition Area - Parma; Safety Assessment of Cosmetics in the EU | Vrije Universiteit Brussel; Medical Devices – EU Directives, Guidance, CE Marking and ISO Standard Certifications | CfPIE; Quality Assurance Systems Based on ISO 9001 and GMP & Auditing and Self-Inspections | European Compliance Academy; Registration of Pharmaceuticals in Europe | IIR/Infoma Life Sciences; e-CTD electronic dossier management for the pharmaceutical industry | Extedo GmbH.

Publications

Over 10 posters or oral communications at scientific or professional meetings.

Interests

Entrepreneurship, business administration and management, project management, regulatory and scientific affairs, product development process. Tennis, golf, soccer, associative work, theatre, agriculture, natural therapies and remedies, etc.

January 2015