Curriculum Vitae

Personal information

Ana Andre

Work experience

1. Employer: INFARMED – Autoridade Nacional do Medicamento e Produtos de Saúde, I.P.
    - Start date: 052008
    - End date:
    - Position: Project Manager _ Scientific Evaluation Unit
    - Activities: Regulatory affairs _ Management of the Pre_ and Post_Authorisation Evaluation of Medicines for Human Use of the Centralised Procedure; collaboration and involvement in the tasks related to the portuguese intervention in the CHMP; Linguistic review and technical assessment of the quality of the Product Information for medicinal products in the Centralised Procedure;
    - Country: Portugal
2. Employer: Community Pharmacy, (Portugal)
    - Start date: 122006
    - End date: 042008
    - Position: Clinical Pharmacist
    - Activities: All community_pharmaceutical activities were covered: Advice to the general public,pharmaceutical care and prescription validation.
    - Country: Portugal

Education and training

1. Subject: Faculty of Pharmacy, University of Lisbon
    - Start date: 092000
    - End date: 102006
    - Qualification: Degree in Pharmaceutical Sciences
    - Organisation: All subjects regarding a Degree in Pharmaceutical Sciences
    - Country: Portugal

Additional information

Publications

Projects

Memberships

Member of the Working Group on Quality Review of Documents QRD _ European Medicines Agency (2011_ Present)

Other Relevant Information

Training sessions, webinars and other: SmPC Advisory Group Webinar – Safety Information II: When does a risk justify a contra_indication, a warning or a precaution for use? _ June 2012 SmPC Advisory Group Webinar – From Scientific assessment to practical information in SmPC and Package Leaflet _ March 2012 SmPC Advisory Group Webinar – SmPC Information on posology and special populations _ November 2011 TOPRA – The organisation for Professionals in Regulatory Affairs _ September 2011 _ CRED LCM Variations _ September 2011 TOPRA – The organisation for Professionals in Regulatory Affairs _ CRED summary of product characteristics _ July 2011 Training in the MedDRA: Coding, Safety Data Analysis and SMQs _ November 2011 INFARMED, I.P. _ Regulatory affairs skills _ gestAR _ 2010 _ present INFARMED, I.P. _ Training session entitled "Quality System Management" _ February 2010 TINFARMED, I.P. _ Training sessions on the readability and braille of the labelling and package leaflets _ January and April 2009 _ Oral presentations under various themes, namely centralized procedure and technical assessment of the product information