



## Curriculum Vitae

Personal information **Patrizia Apollonio**

### Work experience

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1. Employer: AIFA (Italian Medicines Agency)
  - Start date: 042004
  - End date:
  - Position: Pharmacist
  - Activities: Biological Medicinal Products Assessment, Regulatory Affairs, Mutual Recognition Procedure, Decentralised procedure, National procedure, Variation and Renewal procedure
  - Country: Italy
2. Employer: INDENA SPA
  - Start date: 071999
  - End date: 042004
  - Position: Pharmacist
  - Activities: Experimental and preclinical research of the new active substances in chemotherapy for the gynecology tumors Responsible for the Pharmacology and Toxicology Laboratory in vivo
  - Country: Italy
3. Employer: Institute of Pharmacology and Pharmacognosy, Department of Pharmacy, University of Rome
  - Start date: 011998
  - End date: 071999
  - Position: Pharmacist
  - Activities: Experimental thesis in Pharmacology, Pharmacognosy and Toxicology, laboratories of pharmacology, in vivo and in vitro experimental research, and analytical technology (HPLC, Spettrophotometry etc.)
  - Country: Italy

### Education and training

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1. Subject: Department of Obstetrics and Gynecology , Catholic University of the Sacred Heart,
  - Start date: 052000
  - End date: 072004
  - Qualification: Doctorate in Molecular Pathology in gynecology oncology
  - Organisation: Doctorate in Molecular Pathology in gynecology oncology New Taxanes for the treatment of gynecology tumors
  - Country: Italy
2. Subject: Institute of Pharmacology and Pharmacognosy, Department of Pharmacy, University of Rome
  - Start date: 091993
  - End date: 091999
  - Qualification: Degree in Chemistry and Pharmaceutical Technology
  - Organisation: Experimental thesis in Pharmacology, Pharmacognosy and Toxicology, laboratories of pharmacology, in vivo and in vitro experimental research, and analytical technology (HPLC, Spettrophotometry etc.)
  - Country: Italy
3. Subject: Department of Pharmacy and Medicine at the University of Rome
  - Start date: 102014
  - End date: 102016
  - Qualification: Master II level in Regulatory Science Medicines
  - Organisation: Pharmaceutical Regulatory Science: regulatory framework, research and development of the drug, registration of a drug (Dossier), quality control areas, GMP, GLP, GCP, pharmacovigilance inspections, medicinal products for advanced therapies, vaccines and herbal, health economics and drug economy, HTA
  - Country: Italy

### Additional information

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Publications

Projects

Memberships

Other Relevant Information