

Curriculum Vitae

Personal information Yanna-Marina Chevalme

Work experience

- 1. Employer: ANSM
 - Start date: 2001
 - End date:
 - Position: Diagnostic Products (radiopharmaceuticals and contrast media) Assessor
 - Activities: _ marketing authorisation evaluator for radiopharmaceutical (pharmaceutical, preclinical and clinical) _ scientific advice on development and clinical trials for diagnostic agents and therapeutic radiopharmaceuticals. _ Member of the working party on radiopharmaceuticals at EMA (European Medicines Agency). • Rapporteur for the core PIL, core FDG SmPC and core FNA SmPC • Delegate of Afssaps and Ansm at OCDE concerning the global market of technetium_99m
 - Country: France
- 2. Employer: AFSSAPS
 - Start date: 1992 End date: 2001

 - Position: Radiopharmaceuticals Assessor
 Activities: _ marketing authorisation evaluator for radiopharmaceutical (pharmaceutical, preclinical and clinical) _ advice on PSUR. _ Member of the working party on radiopharmaceuticals at Bruxelles then EMEA (European Medicines Agency).
 - Country: France
- 3. Employer: AFSSAPS

 Start date: 1987
 - End date: 1992

 - Position: Radiopharmaceuticals Assessor Activities: _ marketing authorisation dossier on pharmaceutical
 - Country: France

Education and training

1. Subject: University of Pharmacy (Paris XI)

Start date: 1987 End date: 1987 Qualification: PharmD Organisation:

Country: France 2. Subject: University of Law (Sceaux)Start date: 1986

Start date: 1986
End date: 1986

Qualification: Master in Health Law Organisation:

Country: France

Additional information

Publications

Majors Representative references for the past ten years 1. Chevalme Y. How the French Medecine Authority addressed the technetium crisis information. J Nucl Med. 2009; 50 (Supplement 2): 1369 2. Talbot JN, Kerrou K, Montravers F, Nataf V, Chevalme Y. FDOPA PET has clinical utility in brain tumour imaging: a proposal for a revision of the recent EANM guidelines. Eur J Nucl Med Mol Imaging 2007; 34 (7): 1131_2 3. Talbot JN, Chevalme Y. Current evaluation of the clinical utility of Fluoromethylcholine_(18F) PET/CT in prostate cancer. Braz. arch. biol. technol. [online]. 2008; 51 (n.spe): 71_75. 4. Chevalme Y, Montravers F; Vuillez JP, Zanca M, Fallais C, et al..FDOPA_(18F): a PET radiopharmaceutical recently registered for diagnostic use in countries of the European Union. Braz. arch. biol. technol. [online]. 2007; 50 (n.spe): 77_90.5. Chevalme Y. Registration of pharmaceutical and radiopharmaceutical drugs: French and European regulations. Med Nucl 2005; 29 (4): 165_16

Projects

Memberships

Other Relevant Information

_ Quality insurance and normes ISO 9000 _ Since 1996 regular attendance of French congress of nuclear medecine and of radiology, _ Since 2000 regular attendance to the European congress of Nuclear Medicine (EANM) and Radiological Society of North America (RSNA)