



## Curriculum Vitae

Personal information **anissa benlazar**

### Work experience

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1. Employer: ANSM, French Health Authority
  - Start date: 032024
  - End date: -
  - Position: head of oncohematology heematology and nephrology
  - Activities: \_ Ensure functional links and information flow with other technical and medical divisions and the hierarchy, \_ Ensure the organization of work and the management of priorities within the team, \_ Ensure the quality, the respect of deadlines and the consistency of deliverables, \_ EMA centralized procedures referent for oncohematology division.
2. Employer: European Medicines Agency
  - Start date: 062022
  - End date:
  - Position: Oncology European Specialised Expert Community (ESEC\_O) Member
  - Activities:
3. Employer: ANSM, French Health Authority
  - Start date: 082022
  - End date: 122022
  - Position: Interim head of oncohematology heematology and nephrology (maternity leave replacement)
  - Activities: \_ Ensure functional links and information flow with other technical and medical divisions and the hierarchy, \_ Ensure the organization of work and the management of priorities within the team, \_ Ensure the quality, the respect of deadlines and the consistency of deliverables, \_ EMA centralized procedures referent for oncohematology division.
4. Employer: ANSM, French Health Authority
  - Start date: 042021
  - End date:
  - Position: Clinical assessor \_ Oncohematology division
  - Activities: \_ Assessment of safety and efficacy data within Marketing Authorization applications, Clinical Trials applications and early and compassionate access programs, \_ Determination of the Benefit / Risk balance with regards to CT results and therapeutic strategies and alternatives \_ For all assessed procedures and MAH's submitted documents, writing of a critical assessments, \_ Drug shortage anticipation and management with MAHs \_ Expertise at the CHMP, HAS (CT) and the National Cancer Institute (INCa) \_ Ongoing involvement in the ANSM's Permanent Scientific Oncology\_oncohematology Expertise Committee.
5. Employer: ANSM, French Health Authority
  - Start date: 082019
  - End date: 042021
  - Position: Pharmacovigilance assessor \_ Oncohematology division
  - Activities: \_ Scientific and regulatory critical analysis of safety data from clinical trials and post\_marketing experience (MA Applications, safety signals, PSURs, RMPs, PASSs, MA variations and renewals, line and indications extensions and compassionate use programs) \_ For all assessed procedures and MAH's submitted documents, writing of a critical assessments, \_ Expertise at the PRAC.
6. Employer: ANSM, French Health Authority
  - Start date: 022019
  - End date: 082019
  - Position: Pharmacovigilance Assessor Trainee, Oncology division
  - Activities:
    - \_ Safety reports evaluation \_ RMP evaluation \_ Signal management \_ In depth analysis of Hyperprogressive Disease under PD\_(L)1 drugs
7. Employer: Pierre Fabre Oncology
  - Start date: 042018
  - End date: 092018
  - Position: Pharmacovigilance Assessor Trainee, Oncology/Neurology
  - Activities:
    - \_ ICSRs management \_ Causality assessment \_ Follow up of clinical trials' PV activities \_ Literature review
8. Employer: University Hospital
  - Start date: 102016
  - End date: 072017
  - Position: Internship in pharmacy \_ Pharmacology \_ Toxicology \_ Biochemistry \_ Hematology

### Education and training

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1. Subject: Sorbonne University \_ Saint Antoine Hospital
  - Start date: 102021
  - End date: 062022
  - Qualification: Clinical Oncohematology Diploma
  - Organisation: Acquired in addition to the theoretical bases, the practical biological,

cytological, hemostasis and hemobiology bases necessary for the diagnosis and therapeutic management of hematologic malignancies

2. Subject: Pharmacy University, Paris Sud and Paris Descartes
  - Start date: 102018
  - End date: 072019
  - Qualification: Masters degree in Toxicology, Risk Evaluation and Pharmacovigilance
  - Organisation: Objectives of the Masters' degree: \_ Critically analyze scientific articles and regulatory reports to quickly identify relevant information, \_ Use experimental approaches (toxicity evaluation models) for assessing the risks, \_ Master regulatory toxicology environment, \_ Master the essential knowledge of health products' vigilances including regulatory aspects.
3. Subject: Pharmacy University, Paris Sud
  - Start date: 092017
  - End date: 092018
  - Qualification: Masters degree in drug sciences
  - Organisation:
4. Subject: Pharmacy University
  - Start date: 092011
  - End date: 092017
  - Qualification: Pharm\_D
  - Organisation: General pharmacy diploma with a specialization in pharmaceutical industry and toxicology Thesis: Mechanism and biomarkers of hyperprogressive disease during PD\_1/PD\_L1 blockage therapies in patients with solid tumors

## Additional information

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[Publications](#)

[Projects](#)

[Memberships](#)

**Other Relevant Information** \_ C2 English level \_ Volunteer for 6+ years in an association supporting children with cancer