

Curriculum Vitae

Personal information **Meggy Van Nuenen-Cox**

Work experience

1. Employer: GE Healthcare
 - Start date: 052005
 - End date: 072011
 - Position: Quality Officer / Qualified Person
 - Activities: In this role I achieved to set up a standardized complaints process within GE Healthcare international as a member of the global GE Healthcare complaints team. In addition, I introduced a cleanroom behavior monitoring program. I also acted as an auditor internal and external. Furthermore, I have set up a process of annual product reviews and I was a subject matter expert on sterility assurance within GE Healthcare Eindhoven. Finally I was a QP within GE Healthcare Eindhoven and I was accepted as QP for Norway.
 - Country: Netherlands
2. Employer: Dechra
 - Start date: 072011
 - End date: 102015
 - Position: Qualified Person
 - Activities: In this role, I was the sterility expert and subject matter expert during investigations. I was the lead for the update production simulation trail runs to the current guidelines. I have implemented documented evidence of review of batch records. In addition, I was responsible for audit organization, introduction of the war room/ back and the prep room principle. Furthermore, I have gained knowledge regarding different kind of drugs like premix, powders, sterile and aseptic products.
 - Country: Netherlands
3. Employer: Medimmune/Astrazeneca
 - Start date: 112015
 - End date: 082017
 - Position: Manager Fill & Finish
 - Activities: In this position, I am responsible for the operation management of manufacturing activities at the filling department, streamlining of production processes and cost reduction, trouble shooting and life cycle management of manufacturing related activities. This included managing a department of 16 production technicians and 2 shift coaches. I obtained audit experience including FDA, IGZ and third party audits. In addition, I was involved in the implementation of the new filler line a RABS system.
 - Country: Netherlands
4. Employer: Medimmune/AstraZeneca
 - Start date: 082017
 - End date: 102019
 - Position: Production Pharmacist
 - Activities: In this position, I am responsible for the quality of the product produced at Astrazeneca by streamlining and improving of production processes, deviation and CAPA management within manufacturing and reviewing of batch records.
 - Country: Netherlands
5. Employer: IGJ
 - Start date: 102019
 - End date:
 - Position: Coordinating/Specialist Inspector
 - Activities: GMP/GDP inspections (inter)national.
 - Country: Netherlands

Education and training

1. Subject: ECA
 - Start date: 032020
 - End date: 032020
 - Qualification: GMP and Quality for Radiopharmaceuticals
 - Organisation: Quality aspects for producing radiopharmaceuticals
 - Country: Austria
2. Subject: NSF
 - Start date: 022019
 - End date: 022019
 - Qualification: A_Z of Sterile Products Manufacturing
 - Organisation: Detailed training on GMP volume 4, annex 1
 - Country: United Kingdom
3. Subject: University of Utrecht
 - Start date: 091997
 - End date: 042005
 - Qualification: Pharmacist
 - Organisation: Specialization postdoctoral on drugs in the industry including internship at validation department, Solvay pharmaceuticals Olst (7). Specialization courses: pharmaceutical education including 3 months as student assistant (8), entrepreneur (7). Research project: DNA characterization to find the root cause for sickle_cell anemia at the clinical chemical department UMC Utrecht including process development with Polymerase Chain Reaction (PCR).

- Country: Netherlands
- 4. Subject: RADBOUD UNIVERSITY OF NIJMEGEN
 - Start date: 2016
 - End date: 2016
 - Qualification: Radiation Protection expert level
 - Organisation: certification for radiation safety personnel for companies and institutions using radioactive materials according to demands in Dutch legislation ('Kernenergiewetvergunning') and licenses. The level of this course is post_academic.
 - Country: Netherlands
- 5. Subject: NSF
 - Start date: 2006
 - End date: 2006
 - Qualification: Roles and responsibilities of a QP
 - Organisation: In depth knowledge on GMP volume 4 annex 16
 - Country: United Kingdom
- 6. Subject: academie voor toezicht
 - Start date: 112019
 - End date: 062020
 - Qualification: leertraject startende inspecteur
 - Organisation: Oriëntatie op de omgeving: maatschappelijke context van het toezicht
 - Inspectieoptreden: _plannen en voorbereiden _Objectief waarnemen en oordeelsvorming
 - _Objectief interveniëren en communiceren van afspraken _Rapporteren
 - Country: Netherlands

Additional information

Publications None

Projects None

Memberships Member of the Dutch Industrial Pharmacist Association (NIA)

Other Relevant Information