

QA Officer

Job ID

REQ-10072169

Mar 23, 2026

LOC_IT

About the Role

Major Accountabilities:

- Contribute to assuring the validation/qualification status of the production site, equipment, training of personnel and management of quality documentation.
- Responsible for the provisional release for the shipment of batches.
- Work in shift with other QA officers to oversight the production and quality control activities.
- Archiving and support in managing the site GMP documentation.
- Review of batch records and assure the timely closure of the manufactured batches.
- Contribute to maintaining the local quality system as per GMPs and corporate guidelines and in assuring the respect of the GMPs and Health Authorities requirements at local level.
- Support the QP in the preparation of batches release documents.
- Involvement in investigation of deviation, OOS, complaints, CAPA, change control implementation and redaction.
- Collaborate and support during the external audits by the authorities and corporate audits.
- Contribute to redaction and review of SOPs, records, protocols and reports according to GMPs, National/ Corporate Guidelines and health authorities' requirements.

Essential requirements:

- Scientific degree.
- Previous experience in a similar role within a sterile production environment.
- Good knowledge of GMP.
- Available to work in shifts, including night shifts and weekends (on a regular basis)
- Fluent in Italian. Good knowledge of English.

Role Requirements

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Division

DIV_TO

Business Unit

Quality

Location

LOC_IT

Site

Ivrea

Company / Legal Entity

IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl

Functional Area

FCT_QA

Job Type

Full time

Employment Type

Regular

Shift Work

No

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