Call for GMP auditor(s) – 01/2023

About QUAMED

QUAMED is a not-for-profit association with a mission to improve access to quality medicines for all. QUAMED is a global membership organization that brings together international Non-Governmental Organizations (NGOs), pharmaceutical procurement centers, and engaged individuals focused on assuring the quality of medicines delivered in low- and middle-income countries.
As part of its development, QUAMED is looking for independent and experienced auditors who are interested to become validated QUAMED auditors. GMP auditors are professionnals free from conflict of interest specialized in GMP (Good Manufacturing Practices) who will conduct on-site audits of pharmaceutical suppliers. Their task will be to perform assessments of pharmaceutical suppliers using GMP (Good Manufacturing Practices) standards. The aim of these GMP audits is to provide to QUAMED customers an independent and documented feedback of the level of compliance of the audited organization(s) regarding Good Manufacturing Practices. Audits are full assessment (usually 4 or 5 days on site). Assignments will be on a short-term basis (usually one to three weeks). QUAMED has an internal validation process for auditors. For more information, please visit our FAQs on our website : https://quamed.org/audit/

Qualification required

Educational Background
Degree/diploma in pharmacy are preferred.

Qualifications and experiences

- GMP inspectors who are officials (in activity or not in activity for less than 5 years) of stringent regulatory authorities (SRA), WHO-listed authorities (WLA) or international organisations (such as WHO prequalification program (WHO-PQ))
- For other profiles:
    - Good working knowledge of the current WHO GMP guidelines.
    - Suitable qualifications, knowledge, and experience of Quality Assurance in manufacturing of medical products.
    - Recent (less than 5 years) practical experience as GMP auditor, according to international standards (EU/USFDA/WHO).
    - Proven regular training (attestation, certification, etc.) in current standards and knowledge for assessment in pharmaceutical field.
- Low-and-middle income countries professional experiences is an asset

Others skills

- Ability to work in English is required
- Ability to work also in French or Spanish or Arabic is an asset

- Communication and reporting skills

How to apply

- Applications (Curriculum vitae including a short presentation describing your abilities to meet the essential requirements (max 1500 words) as well as names and contact details of a minimum of 2 references) should be sent by email to l.lavergne@quamed.org
- All the applicants will be informed by e-mail of the results of the selection process.
- The final deadline for application is the 28/02/2023

Call for GSDP auditor(s) – 01/2023

About QUAMED

QUAMED is a not-for-profit association with a mission to improve access to quality medicines for all. QUAMED is a global membership organization that brings together international Non-Governmental Organizations (NGOs), pharmaceutical procurement centers, and engaged individuals focused on assuring the quality of medicines delivered in low- and middle-income countries.

As part of its development, QUAMED is looking for independent and experienced auditors who are interested to become validated QUAMED auditors. GSDP auditors are professionnals free from conflict of interest specialized in GSDP (Good Storage and Distribution Practices) who will conduct on-site audits of pharmaceutical suppliers. Their task will be to perform assessments of pharmaceutical suppliers using World Health Organization (WHO) GSDP (Good Storage and Distribution Practices) standards. The aim of these GDSP audits is to provide to QUAMED customers an independent and documented feedback of the level of compliance of the audited organization(s) regarding Good Distribution and Storage Practices. Audits are full assessment (usually 2 or 3 days on site) and can be part of a Local Market Assessment (LMA) including several pharmaceutical distributors. Assignments will be on a short-term basis (usually one to three weeks). QUAMED has an internal validation process for auditors. For more information, please visit our FAQs on our website : https://quamed.org/audit/

Auditors are professionnals free from conflict of interest specialized in GSDP who will conduct on-site audits of pharmaceutical suppliers. On-site audit visits usually last between two to five days.

Qualification required

Educational Background

Degree/diploma in pharmacy is preferred.

Qualifications and experiences

- GSDP/GMP inspectors who are officials (active or less than 5 years retired) of stringent regulatory authorities (SRA), WHO-listed authorities (WLA) or international organisations (such as WHO prequalification program (WHO-PQ))
- For other profiles:
    - Good working knowledge of the current WHO GSDP guidelines.
    - Suitable qualifications, knowledge, and experience of Quality Assurance in procurement and supply management of medical products.
    - Recent (less than 5 years) practical experience as GSDP auditor, according to international standards (EU/USFDA/WHO).

  - Proven regular training (attestation, certification, etc.) in current standards and knowledge for assessment in pharmaceutical field.
- Low-and-middle income countries professional experiences is an asset

Others skills

- Ability to work in English is required
- Ability to work also in French or Spanish or Arabic is an asset
- Communication and reporting skills.
- Ability to work with Word and Excel as well as familiarity with cloud-based tools

How to apply

- Applications (Curriculum vitae including a short presentation describing your abilities to meet the essential requirements (max 1500 words) as well as names and contact details of at least 2 references) should be sent by email to l.lavergne@quamed.org
- All the applicants will be informed by e-mail of the results of the selection process.
- The final deadline for application is the 28/02/2023

Call for GSDP expert(s) – 01/2023

About QUAMED

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As part of its development, QUAMED is looking for independent and experienced experts who are interested to become validated QUAMED experts. Their task will be to perform assessments of pharmaceutical suppliers using World Health Organization (WHO) GSDP (Good Storage and Distribution Practices) standards.

Experts are free from conflict of interest professionals who will conduct GSDP technical visits, a short assessment of distributors conducted on-site or remotely. The aim of the GDSP technical visits is to provide QUAMED customers with an independent, rated and documented feedback of the level of compliance of the visited organization(s) with GSDP standards. Technical visits are part of a Local Market Assessment (LMA) including several pharmaceutical distributors. Assignments will be on a short-term basis (usually one to three weeks). QUAMED has an internal validation process for experts. For more information, please visit our FAQs on our website : https://quamed.org/audit/

Qualification required

Educational Background
Degree/diploma in pharmacy is preferred.

Qualifications and experiences

- Good working knowledge of the current WHO GSDP guidelines.
- Suitable qualifications, knowledge, and experience in quality assurance in procurement and supply management of medical products.
- Recent (less than 5 years) practical experience as GSDP expert, according to international standards (EU/USFDA/WHO).
- Proven regular training (attestation, certification, etc.) in current standards and knowledge for assessment in pharmaceutical field.
- Low-and-middle income countries professional experiences is an asset

Others skills

- Ability to work in English is required
- Ability to work in French, Spanish or Arabic is an asset
- Communication and reporting skills
- Ability to work with Word and Excel as well as familiarity with cloud-based tools

How to apply

- Applications (Curriculum vitae including a short presentation describing your abilities to meet the essential requirements (max 1500 words) as well as names and contact details of at least 2 references) should be sent by email to l.lavergne@quamed.org
- All the applicants will be informed by e-mail of the results of the selection process.
- The final deadline for application is the 28/02/2023

Call for MQAS auditor(s) – 01/2023

About QUAMED

QUAMED is a not-for-profit association with a mission to improve access to quality medicines for all. QUAMED is a global membership organization that brings together international Non-Governmental Organizations (NGOs), pharmaceutical procurement centers, and engaged individuals focused on assuring the quality of medicines delivered in low- and middle-income countries.

As part of its development, QUAMED is looking for independent and experienced auditors who are interested to become validated QUAMED auditors. MQAS auditors are professionnals free from conflict of interest specialized in MQAS (Model Quality Assurance System for Supply Agencies) and GSDP (Good Storage and Distribution Practices) who will conduct on-site audits of pharmaceutical suppliers. Their task will be to perform assessments of pharmaceutical suppliers using MQAS (Model Quality Assurance System for Supply Agencies) and World Health Organization (WHO) GSDP (Good Storage and Distribution Practices) standards. The aim of these MQAS audits is to provide to QUAMED customers an independent and documented feedback of the level of compliance of the audited organization(s) regarding MQAS. Audits are full assessment (usually 3 days on site) and can be part of a Local Market Assessment (LMA) including several pharmaceutical distributors. Assignments will be on a short-term basis (usually one to three weeks). QUAMED has an internal validation process for auditors. For more information, please visit our FAQs on our website : https://quamed.org/audit/

Qualification required

Educational Background

Degree/diploma in pharmacy are preferred.

Qualifications and experiences

- GSDP/GMP inspectors who are officials (in activity or not in activity for less than 5 years) of stringent regulatory authorities (SRA), WHO-listed authorities (WLA) or international organisations (such as WHO prequalification program (WHO-PQ))
- For other profiles:
    - Good working knowledge of the current WHO GSDP and WHO MQAS guidelines.
    - Suitable qualifications, knowledge, and experience of Quality Assurance in procurement and supply management of medical products.
    - Recent (less than 5 years) practical experience as MQAS auditor or GSDP auditor, according to international standards (EU/USFDA/WHO).
    - Proven regular training (attestation, certification, etc.) in current standards and knowledge for assessment in pharmaceutical field.
- Low-and-middle income countries professional experiences is an asset

Others skills

- Ability to work in English is required
- Ability to work also in French or Spanish or Arabic is an asset
- Communication and reporting skills

How to apply

- Applications (Curriculum vitae including a short presentation describing your abilities to meet the essential requirements (max 1500 words) as well as names and contact details of a minimum of 2 references) should be sent by email to l.lavergne@quamed.org
- All the applicants will be informed by e-mail of the results of the selection process.
- The final deadline for application is the 28/02/2023