QUAMED services

Our services

- MQAS Audit
- GSDP Audit
- **Technical visit GSDP**
- GMP Audit
- GMP technical visit
- **Evaluation of sources**
- **Evaluation of product files**

- Evaluate the quality of products offered by a manufacturer based on their FPP* product files (PD)

  Finished *Pharmaceutical Product*
- 4 weeks from receipt of RFP (via client)
- 2 qualified and validated QUAMED evaluators chosen according to the expertise required for the evaluation of the FPP file.
- Collection of product information (Name of manufacturer, Address, Country of production, ...) and evaluation of the product file (GMP status, API specifications, Finished product specifications, Stability studies, Packaging/label/notice, Safety and efficacy)
- QUAMED Guideline "Product Dossier Evaluation" + Interagency FPP Questionnaire (IAPQ) + QUAPAS QUAMED

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- **Report**:
  - Transmission of the report 4 weeks after receipt of the RFP
  - Any changes to the submitted RFP must be reported to QUAMED within 3 years of the assessment
  - Active follow-up by QUAMED (if client interest is confirmed)
  - Deposit on the QUAMED database (valid for 3 years).