Auditor for the Analytical R&D and QC unit

At Protalix, our focus is to develop, produce and commercialize potentially better therapies for a variety of illnesses through cutting-edge research and development and our proprietary plant cell-based protein expression platform.
If you wish to be a part of our success and join a dynamic and professional team that allows personal and professional growth, we are looking for you!

Responsibilities

- Evaluate the adequacy and compliance of systems, operations, and practices against regulations and company documentation
- Ensure that laboratory operations are completed according to established Standard Operating Procedures (SOPs)
- Check and review data in compliance with data Integrity requirements
- Draft and control SOPs, worksheets and other Quality Control documentation
- Involvement in analytical method validation and qualification studies
- Management of specifications for raw material, IPC, DS and DP
- Work with pharmacopeias in order to audit the testing of different materials

Requirements

- Preferred: An understanding of GMP (Good Manufacture Practice) regulatory guidelines.
- Preferred: Knowledge in an auditing process in a GMP environment
- Essential: Computing skills, including use of spreadsheets (e.g. Word, Excel)
- Quality orientation and high attention to detail
- Work in an orderly and systematic approach
- Ability to work collaboratively as well as independently, with deferent QC members
- Self-motivated & strong interpersonal and communication skills
- Good writing and verbal communication skills, both in English and Hebrew language
- Analytical capabilities

Please send your application to: hr-cv@protalix.com