We go beyond the obvious
and make a positive impact on people's lives

Serialization Specialist
(Ref: SS1123)

RAFARM is an innovation-driven, dynamically growing pharmaceutical company and a well-established European manufacturer with an outward-looking orientation that invests 13% of net turnover in Research and Development. We introduce high technology in our state-of-the-art sterile manufacturing plant and create new production lines with cutting-edge technology and robotic equipment. RAFARM is a place where our people evolve and are full of passion and interests. We are a dynamic team that empowers talent, embraces diversity and accelerates development.

The role main accountabilities will be the following:

- Management of business partners and ensuring interfaces between systems is maintained and validated according to Rafarm policies and procedures
- Championing CMO onboarding activities as it relates to serialization
- Creating/Editing Level 1 Print files: Data Matrix Configuration, Human Readable Lines Configuration. Managing Level 2 and Level 3 software between batches to ensure smooth operations. Managing & maintaining Level 4 software to ensure uninterrupted supply chain operation
- Coordinate all serialization activity and system interfaces with the appropriate team such as: IT, Production, Quality and Regulatory
- Manage change controls and deviations associated with Serialization as well as issuing CAPAs as required
- Manage Serialization Issue Log and be able to interpret EPCIS events according to the GS1 standards
- Lead resolution of all issues associated with serialization and provide department SOPs
- Audit trail and data review with the understanding of 21 CFR PART 11.
- Development of Design Qualifications, Functional Requirements, User Requirements, Unit test scripts, UAT scripts, validation plans, validation summary report, traceability matrix, IQ/OQ/PQ
- Contribute in the definition of the future technology roadmap by identifying value adding technology opportunities, and support the implementation of SAP S/4HANA

Experience & Qualification Standards:

- BSc in Computer science, Engineering, Economics or relevant discipline
- A at least 2 years' experience in the domain of serialization in the pharmaceutical sector
- Proven experience in Systech's UniSeries and UniTrace serialization platform
- Strong knowledge of Global Serialization Standards (GS1), regulatory requirements and guidance
- Proficiency in SAP (MM & PP) will be a plus
- Enthusiastic and able to take ownership of problems and provide solutions
- Excellent oral communication & report, business correspondence & procedure-writing skills both in Greek and English languages
- Highly detail-oriented and organized, able to work both independently and as a team player with a positive attitude

The Company offers: Competitive remuneration package along with challenging career development opportunities within an innovation orientated organization

The candidates with the above qualifications can send their CVs to: hrdep@rafarm.gr