Job Title: Formulation Scientist

Location: Bridgewater, NJ

Solaris Pharma Corporation is a leading generic corporation with the vision of improving the patient's quality of life through development of niche pharmaceutical products. Solaris Pharma Corporation has a fully equipped Research & Development facility with expertise in the development of generic and specialty products. Solaris Pharma Corporation develops specialty dermatology products that have a relatively high barrier to entry due to manufacturing complexities and high-end clinical patient study demands. With a thorough knowledge of the development life cycle and well-trained, committed scientific personnel, Solaris Pharma Corporation has streamlined processes that are efficient in reducing development costs and shortening timelines, without compromise in quality.

Solaris Pharma Corporation is currently engaged in the development of several generic dermatology products and is committed to establish itself as a pacesetter in dermatology pharmaceuticals.
Solaris Pharma Corporation, is looking for a full time, Associate Scientist.

Responsibilities:

Independently creating and implementing formulation development strategies. Design, direct and/or conduct pre-formulation, formulation development, process development, process optimization, scale-up and clinical manufacturing on assigned projects of **Generic Formulations.**

- Prepare and execute the lab scale developmental batches for Topical products (Creams, Ointments, Gels etc...)
- Formulation and interpretation of results.
- Writing lab note book and compilation of results for each project in up-to- date fashion.
- Conduct a pre-formulation studies and stability of development batches as and when required during development.
- Review and authorize reports/documents such as master formulas and other key reports/ documents.
- Conduct IVRT/IVPT experiments as per guidance from Lead Scientist
- Help lead scientist for Development of Rheological characteristic for Semi-Solid Products
- Assist in the exhibition of ANDA submission batch and execution of protocols for Topical Products
- Coordinate with contract research organizations for scale up & clinical batches and provide technical assessments
- Help lead Scientist in daily routine formulation activity

Qualifications:

- BS, with at least 2-3 years' experience Or Master with 0-1 year in Pharmaceutics or related field in semi-solid (preferred) or related product development
- Must have excellent communication skills, organizational ability and good documentation capabilities
- Working experience in the development of semi-solid generic products is a plus.
- Experience with dosage forms such as suspensions and emulsions is a plus
- Excellent technical writing skills (Feasibility reports on new opportunities, technical due diligence supportive documents, experimental protocols, development reports, and technology transfer reports)
- Experience in both brand and generic industry is a plus