BlueReg Group is a consultancy company specialized in Development, Regulatory Affairs, quality management and Pharmacovigilance for the Pharmaceutical Industry.
Based in Paris, Sophia Antipolis and London, BlueReg Group has over 100 clients ranging from large multinationals to small start-up companies, from innovators to generic companies.

We are recruiting:
Senior Consultant, Regulatory Affairs (EU and UK/Ireland)

Your key duties and responsibilities:

- Is a source of regulatory expertise in the development, registration and post-licensing activities of pharmaceutical products
- Project management activities
- People management / Review of work of junior personnel and BlueReg partners
- Ensures high quality and on time delivery of consulting services to clients.
- Contributes to the effective functioning and to the business growth of BlueReg EU

Your profile :

- Pharmacist, Life Sciences Graduate, ideally with post-graduate qualification.
- Sound knowledge of pharmaceutical regulations and guidelines.
- Significant regulatory affairs experience including a successful track record in the registration of medicinal products
- A least 10 year experience in the pharmaceutical industry, broadly based Regulatory Affairs experience, including a successful track record in the registration and maintenance of pharmaceutical products within Europe and UK/Ireland.

You have excellent written and verbal communication skills, good organisational and analytical skills. You are fluent in written and spoken English.

This position is based in High Wycombe, UK

If you are interested in, please send a copy of your CV and a cover letter to contact@blue-reg.com.