Bioscript®
REGULATORY

Senior/Principal Regulatory Writer

Are you an experienced Regulatory Writer working at Senior or Principal Writer level? Are you committed to high standards of delivery and technical excellence? If so, we can offer you new opportunities in a forward-thinking, friendly and dynamic agency.

Location: Alderley Park, East Cheshire

Job type: Permanent; full-time is preferred, although part-time arrangements can be discussed

Benefits: Competitive salary, profit-related bonus, pension, health insurance, 25 days' holiday, and more.

About Bioscript Regulatory
Bioscript Regulatory is part of Bioscript Group, a full-service, global healthcare communications agency delivering communication planning, publications, medical meetings, digital programmes, health economic and regulatory services to pharmaceutical companies. We have grown steadily over the past 12 years, adding new clients and talented new staff every year; we are proud that the clients we worked with at our inception in 2005 continue to work with us.

Duties and responsibilities of the role
As a Senior Regulatory Writer, you will prepare clinical regulatory documents in a diverse range of therapy areas for our pharmaceutical clients, utilising and broadening your existing experience. You will be familiar with documents such as clinical study reports and the common technical document (CTD), and will be able to evaluate scientific data objectively, in order to report results and scientific concepts in a concise and accurate manner. You will also have involvement in client services and will be able to contribute to the strategic planning of regulatory submissions.

In addition to the above, as a Principal Regulatory Writer you will be able to lead the planning and delivery of regulatory submissions and other large regulatory projects.

This position will be based in our offices at Alderley Park, East Cheshire, although we are also willing to consider flexible working arrangements.

Requirements
We are interested to hear from individuals with drug development experience, relevant to the field of regulatory writing. You will need a strong scientific background, including a minimum of a BSc (or equivalent) in an appropriate scientific discipline. You must have excellent written and spoken English, and a good working understanding of clinical development and ICH guidelines for regulatory documents. Successful candidates will have a proactive approach, well-developed project management skills, and confident and effective interpersonal skills. Flexibility for occasional travel within Europe and the United States is desirable.

Interested?
Send your CV and a brief covering letter to careers@bioscriptgroup.com

All offers of employment are subject to candidates' ability to provide suitable documentary evidence of their right to work in the UK.