

Position Available: Senior Regulatory Affairs Project
Manager/Associate Director of Regulatory Affairs
ERA Consulting und Pharmaberatungs GmbH
Our Ref: ERA/DE/APP/42 Date: 7th December 2004



POSITION AVAILABLE

Senior Regulatory Affairs Project Manager/Associate Director of Regulatory Affairs

With office locations in London (UK), near Hanover (Germany), Washington DC (USA), and Brisbane (Australia), a highly experienced team and a proven track record of success, the ERA Group is one of the foremost consulting organisations dedicated to the development of biotechnology-derived medicinal products and other biologics. Alongside product development consulting, we also offer high-level strategic regulatory affairs consulting services in this field. Our clients range from smaller biotech start-ups to the largest international pharmaceutical companies.

The successful candidate for this position will have the following qualifications, experience and skills:

- A degree and higher degree (Masters or Ph.D.) in a relevant subject (e.g. cell biology, biochemistry)
- Experience and a strong track record in biotech product development and/or regulatory affairs
- Proven management skills and strong motivation for teamwork
- Good writing skills (English) and computer literacy (word-processing, databanks etc.)

After an appropriate period of initiation into the company's projects, it is expected that the candidate will assume the position of Associate Director of Regulatory Affairs, reporting directly to the Group Director, assuming overall responsibility for ERA's activities in the German-speaking regions and heading our Walsrode team.

APPLICATIONS:

Applications should be addressed to:

Mrs Sabine Husmann-Holloway
Group Chairperson and Chief Executive Officer
Email: info@eraconsulting.com

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