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A global healthcare leader, Novartis has one of the most exciting product pipelines in the industry today. A pipeline of innovative medicines brought to life by diverse, talented, performance-driven people. All of which makes us one of the most rewarding employers in our field.

Senior Medical Writer / Medical Writer

Horsham, West Sussex

You will write and/or edit the production of high quality clinical documentation for submission to regulatory authorities in support of marketing applications. This will involve you acting as a member of clinical trial teams, participating in the planning of analysis and data presentation to be used for study reports and acting as a documentation consultant to ensure compliance with internal company standards and external regulatory guidelines.

You will also have the opportunity to develop your career as a medical writer, writing and editing

clinical CTD summary documents, co-ordinating medical writing resources for specific clinical programs, working with external vendors and helping to organise the medical writing teams for regulatory submissions.

With a minimum of a life science degree (or equivalent), you will have a knowledge of clinical development and regulatory documentation and an understanding of statistics and data interpretation. You should be a team player with strong interpersonal and presentation skills, and solid written English. A background in medical writing, including experience of managing external medical writers or regulatory knowledge would also be beneficial.

In return you can expect a competitive package, flexible working, flexible benefits scheme (launching January 2009) and generous relocation assistance.

For more details about these and other opportunities, please visit www.novartis.com quoting the reference 44730BR.



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