



**Clinical Safety Manager / Scientist - SERM Pharmacovigilance
Weybridge - Surrey or Stockley Park - West London, South East England**

This is an exciting opportunity to join GSK as a full-time Safety Manager primarily responsible for the continued safety assessment, evaluation and risk management of GSK products or product groups in clinical development and/or the post-marketing setting within the Safety Evaluation and Risk Management (SERM) group.

The Safety Manager ensures that adverse event and other safety information is efficiently evaluated and that safety reports are completed accurately and in a timely manner to meet global compliance and regulatory requirements, identify the need for product labelling updates and drive proactive implementation of risk management initiatives

In this role, you would be responsible for leading risk management activities, providing significant and proactive clinical safety input on cross-functional teams and ensures that appropriate safety objectives and risk minimisation strategies are included in clinical development programs

The ideal candidate will:

- Function independently in cross-functional teams, exhibit a proactive approach, and who is able to lead the safety group within the project teams, with influencing skills and personal strength and impact.
- Understand the big picture as well as having an eye for detail.
- Demonstrate initiative in solving problems, with the ability to manage tasks and projects and be able to prioritise and deliver within timelines.
- Be able to communicate at all levels within a matrix organisation.

In addition, you will play a pivotal role in the writing and creation of regulatory documents, such as PSURs, responses to regulatory enquiries and global core labelling. Working as part of a team, you'll be given full support and training as well as the excellent package you can expect from GSK.

Basic qualifications required:

- Biomedical degree (or equivalent), or higher.

Preferred qualifications:

- Clinical Safety and Pharmacovigilance experience or other relevant experience such as project management within pharmaceutical industry, medical writing, regulatory affairs or clinical development experience.
- Excellent oral and written communication skills.
- Sound computer skills.
- Awareness of international pharmacovigilance requirements for example ICH. Vol 9 Notice to Applicants, and CIOMS initiatives.

Closing date for applications: 31 January 2010.

Please indicate which location (Weybridge or Stockley Park, West London) you are applying for.

To Apply:

Please apply online via GSK's online application system by following the link below and searching for Req. ID 57074.

www.gsk.com/careers/uk-saa-jobsearch.htm

GSK is an equal opportunities employer and is proud to promote an open culture, encouraging people to be themselves and giving their ideas a chance to flourish. To enable GSK to meet its commitment as a two ticks employer please let us know if you have a disability.



Together we can make life better.