

IMPROVING PATIENTS' LIVES THROUGH PASSION.

Director (Physician), Safety Evaluation Risk Management (SERM) – Stockley Park, Uxbridge

At GSK, our evolving pipeline has resulted in the expansion of the Safety Evaluation Risk Management (SERM) team, offering opportunities for Physicians to join this team and take on senior, leading and highly visible roles. Being part of GSK's Global Safety & Pharmacovigilance department, you will lead safety strategies to add to, and continue the development of our highly innovative pipeline.

These key roles offer proactive management of the safety of products at all stages of the life-cycle, and provide a senior point of accountability for the safety of products in clinical development programmes and in the market place. In addition to breaking the mould on how we can develop and deliver medicines to our patients, GSK is leading the way in the development of state of the art tools within the pharmacovigilance arena; this is a safety evaluation and risk management role rather than processing individual adverse event reports.

In this role you will be accountable for delivering the safety strategy, and execution of plans laid out here including risk management activities.

You will:

- lead the discussion and management of safety issues through matrix teams across the global organisation and present your findings to high level clinical teams and GSK
- lead plans for safety data collection, data analysis and interpretation and ensure optimal decisions are made as early as possible; and
- act as an ambassador for GSK in representing safety within the external environment including interactions with regulatory authorities, licensing partners, Independent Data Monitoring Committees/Data Safety Monitoring Boards and external expert groups.

Ideally, you will have proven leadership skills with the ability to influence with credibility both externally as well as at all levels within a matrix organisation. What's more, you will be a highly analytical thinker, having the ability to lead analysis of clinical safety and scientific data, whilst applying sound medical judgment to recognise key issues and provide practical solutions. You will also have highly-effective communication skills and the ability to assimilate and accurately present complex data/information to all levels of the organisation and externally to GSK.

With a demonstrable track record of good decision making and problem resolution based on all relevant information, you will also have the ability to work effectively in an environment characterised by tight timelines and changing priorities.

Self-motivation, and with the intellectual flexibility to continually develop and learn new skills are also critical.

Basic qualifications required;

Medical degree, MD, MBBS or equivalent, with subsequent practical experience of clinical medicine. You will also have an in-depth background in drug safety or other relevant field such as clinical development or medical affairs. Preferred qualifications are GMC-recognised.

For further information and to apply online, please visit our website at www.gsk.com/careers searching for Req ID 68363.

Closing date: Friday 16th March 2012.

As a result of the flow of products through our pipeline, we have vacancies across our therapy areas and would welcome applications from Safety Physicians and Scientists looking for their next career move.

GSK is proud to promote an open culture, encouraging people to be themselves and giving ideas a chance to flourish. We are an equal opportunities employer and welcome applications regardless of gender, age, ethnicity, disability, sexual orientation, country of origin or country of application.

Together we can make life better.



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