Corrective and Preventative Action (CAPA) Manager - Quality

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At Roche, 88,500 people across 150 countries are pushing back the frontiers of healthcare. Working together, we’ve become one of the world’s leading research-focused healthcare groups. Our success is built on innovation, curiosity and diversity. Roche is an equal opportunity employer and strictly prohibits unlawful discrimination based upon an individual’s race, color, religion, gender, sexual orientation, gender identity/expression, national origin/ancestry, mental/physical disability, medical condition, marital status, veteran status, or any other characteristic protected by law.

The Position
Leading and supporting the investigation, root cause analysis and impact assessment of deviations you will contribute to the continuous improvement strategy at Roche. Continuous improvement is an integral part of the Quality Management System (QMS) and you will focus on developing Corrective and Preventative Action (CAPA) plans with affiliates, functions and global process owners to resolve and prevent deviations and improve processes. You will also support the CAPA process specialist in assessing CAPAs, managing process metrics and enhancing the global CAPA process. Establishing strong partnerships with business stakeholders and providing expert guidance relating to CAPA processes, you will develop support that contributes to organisational objectives whilst maintaining a focus on accurate information and reporting. You’ll be proactive in maintaining an up to date knowledge of regulatory requirements and work to ensure that internal policies and external legislation are closely adhered to. You will also deliver regular reports to Senior Management on deviations and the effectiveness of the CAPA process.

Qualifications
With a degree or equivalent in a scientific or quality related field, you will need a significant understanding of quality assurance processes within a pharmaceutical development environment, able to demonstrate applied knowledge of good clinical practice and drug safety alongside an analytical and organised approach. It’s important that you can work effectively and influence others working with many departments and internal or external stakeholders in the context of a large global organisation with experience and ability in project management and team leadership. In return we offer a competitive salary plus the excellent benefits you would expect from a bluechip organisation, including a genuine interest in your development.

Closing date: 13th October 2015
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