



# AFRIKA KOMMT! 2022-2024

An Initiative of German Industry for Future Leaders from Africa

## Candidate Profile: Global Patient Safety Regions Project Specialist (Pharmacovigilance)

<b>Company:</b>	<b>Merck Healthcare KGaA</b>	
<b>Education:</b>	A thorough background in life-sciences, ideally pharmacist. Chemists, Biochemists, Biologists, Neuroscientists etc., and Physicians are also eligible.	
<b>Professional Experience:</b>	3-5 years of experience in the Health Care industry, preferable in post-marketing Pharmacovigilance or Clinical Trials under GCP, Quality Assurance, or Project Management within a multinational organization. Experience of working in a regulatory authority is an extremely valuable asset.	
<b>Additional Qualifications:</b>	<ul style="list-style-type: none"> <li>• Good understanding of global and Regional and local Pharmacovigilance Regulation and Guidance documents</li> <li>• Pragmatic and collaborative approach of implementing regulatory guidance into own area of work</li> <li>• Excellent communication (written and spoken) in English, additional languages are an asset</li> <li>• Initial project management experience, including stakeholder and interface management</li> <li>• Good time management and self-organizational skills</li> <li>• Flexibility and agile approach, ability to make decisions based on limited information</li> <li>• Attention to detail and focus on quality</li> <li>• Ability and willingness to give and accept feedback constructively</li> </ul>	
<b>Division / Department, Place:</b>	Merck R&D Healthcare business, Global Patient Safety (GPS) Germany, Darmstadt. This opening is in GPS Regions, a department acting as bridge between headquarters and affiliates.	
<b>Assignment / Area of Activity:</b>	<ul style="list-style-type: none"> <li>• Project Assignment in Global Patient Safety Regions</li> <li>• Support in Regions improvement projects and strategic initiatives</li> <li>• Interact with R&amp;D functions on collaboration and improvement projects</li> <li>• Contribution to Pharmacovigilance Safety Intelligence regulatory oversight on worldwide changing PV requirements and their impact of the company's PV processes</li> <li>• Support in writing and updating Quality Documents</li> <li>• Support in PV Training activities and education</li> <li>• Support in CAPA Management with Audits and Inspections</li> <li>• Contribute to pharmacovigilance initiatives and projects in Africa</li> </ul>	
<b>Remarks:</b>	The candidate will be based in Germany.	
<b>Preferred Nationality:</b>	All nationalities welcome.	