AFRIKA KOMMT! 2025-2027

An Initiative of German Industry for Future Leaders from Africa

Overall remarks: AK is a fellowship position I.e Alongside making meaningful contributions, the primary focus is on your professional development and learning.

Profile Name: Quality Specialist (f/m/d) at a global Quality Management Department for non-active Medical Devices for one world-wide Manufacturer of Medical Devices and Medicinal Products

Company: **ID: AK14_BB_01**	**B. Braun Melsungen AG / Medical Devices and Medicinal Products**	B BRAUN SHARING EXPERTISE

Education:	Bachelor's degree in medical technology, engineering (or sciences) or comparable degree; a master's degree would be an advantage
Professional Experience:	✓ Work experience in the field of quality management ✓ Ideally knowledge of ISO 13485 ✓ Basic understanding of regulatory requirements and product safety ✓ Experience in conducting trainings; alternatively strong communication and presentation skills ✓ Work experience in the field of project management

Maximum years' experience needed:	Graduate / work experience 1-4y	Other (explain the reason and specify)
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Additional Qualifications:	Experience in: ✓ Good English language skills ✓ Strong communication and presentation skills ✓ Basic microbiological knowledge (may be an advantage) ✓ French language skills (may be an advantage)
Country Focus:	• All nationalities • For information only: We have a Medical Device manufacturing site in Knysna (South Africa) and further affiliates in Kenya (Nairobi), Ghana (Accra), South Africa (Johannesburg), Zambia (Lusaka) and Zimbabwe (Harare).
Division / Location / Department:	Global Quality Management Department in the Headquarter in Melsungen (Germany). Further assignments may be possible at the local quality management department of the Plant Medical Melsungen (Germany) and Almo (subsidiary of B. Braun in Bad Arolsen - Germany).
Planned Tasks & Activities:	Potential areas of assignment and support could be: ✓ Conducting assessments as part of deviation management in the international production environment, including support in the further development of the software applied ✓ Execution of assessments of corrective and preventive measures in the international production environment, including support in the further development of the software applied ✓ Performing assessments as part of change management in the international production environment ✓ Compiling KPI reports and evaluating them in the international production environment ✓ Performing assessments of design verification reports from various development activities

	✓ Performing assessments of qualification and validation activities in the international production environment (incl. clean rooms and sterilization processes) ✓ Quality management activities in the business-to-business environment
Remarks:	Further assignments may be possible at the local quality management department of the Plant Medical Melsungen (Germany) and Almo (subsidiary of B. Braun in Bad Arolsen - Germany) approximately for 1 month