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: Clinical Data Management Fellow

Company ID: **ID:AK_14_ BNT_03**	**BioNTech / Biotech**	Company Logo

Education:	**University degree in Natural Sciences or Life sciences**
Professional Experience:	• Minimum of 4 years of experience in Clinical Data Management (CDM) with good practice in CDM functional activities. • Has good understanding of what basic deliverables CDM is responsible for in clinical studies • Good knowledge of international guidelines (ICH-GCP) for the conduct of clinical research projects. • Experience with at least one Clinical Data Management/Electronic Data Capture System

Maximum years' experience needed:	Graduate / work experience 1-4y	Other (explain the reason and specify)
	☐	Minimum of 4years experience

Additional Qualifications:	• Good communication skills • Detail-oriented • Good problem-solving skills • Forward thinker • Experience in usual software (Word, Excel, Power Point) • Proficiency in English (written and spoken).
Country Focus:	All nationalities from sub-Saharan Africa are welcome to apply. Nationals from the following countries are preferred: Rwanda, Mozambique, Tanzania, Uganda, Gabon and Democratic Republic of Congo.
Division / Location / Department:	Global Clinical Development Operations located in Mainz, Germany .
Planned Tasks & Activities:	• Supports oversight of CRO clinical data management activities guidance from a Senior Manager Clinical Data Manager or an Associate Director Clinical Data Management and in accordance with BioNTech's standards trial oversight process. • Co-creates and/or reviews CRO's data management documents (e.g. Data management plan, CRF, validation plan, etc.) requiring Sponsor's input.

	• Collaborates with CRO's DM to identify requirements for the development and amendments of the clinical database in cooperation with the trial team (Managers Clinical Trial, Medical / Clinical Development experts). • Performs Sponsor User Acceptance Testing of CDM deliverables. • Drive data cleaning activities across functions and ensure CRO is fully aligned with scope and timing • Performs data reviews and quality checks per BioNTech's trial oversight process. • Perform other assigned tasks from CDM management.
Remarks:	At the end of the fellowship to have developed competencies in CDM technical expertise, ability to work cross-functionally and acquired knowledge on how to oversee a CRO for end-to-end CDM activities.