AGNES SITTA KIJO

Mrs. Agnes Sitta Kijo, is a pharmacist, with a Master of Science (MSc) in Pharmaceutical Technology. She has over 23 years of experience in regulation of medicines and medical devices including in vitro diagnostics.

As a Technical Officer, at the Facilitated Products Introduction (FPI) Team in WHO-HQ, Geneva, Switzerland, she is responsible for the Collaborative Registration Procedure for in vitro diagnostics and other facilitated regulatory pathways activities. She also provides technical support related to regulation of medical devices globally and serves as member of the AMRH Joint Secretariat through the African Medical Devices Forum (AMDF) which oversees matters related to regulation of medical devices in Africa.

Before joining WHO in March 2020, she worked as regulator for 19 years and held different technical, advisory and leadership roles which allows her to provide sound and relevant guidance/advice to regulators and regional harmonization initiatives globally.