International Alliance for
Biological Standardization

Maintaining the Quality of Vaccines Through the Use of Standards: Current Challenges and Future Opportunities

Library and Archives Canada, Ottawa, Canada

June 21-22, 2023

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Dr Wim Van Molle is working at Sciensano, Belgium, in the division of Quality of Vaccines and Blood Products, where he is responsible for vaccine batch release and evaluation of the quality part of registration and variation dossiers of vaccines and biologicals. Wim is involved in scientific advice procedures at national and European level and participates as product expert in GMP inspections on the request of the Belgian (FAMHP) and European (EMA) medicines agency. Dr. Van Molle is actively involved in European IMI projects.

At Sciensano, he is manager of the unit for Cellular and Molecular Biology and Biochemistry, involved in quality release testing of vaccines and blood derived medicinal products.

Wim works as auditor for the European Directorate for the Quality of Medicines & Healthcare (EDQM), Strasbourg, France where he is frequently asked to participate in mutual joined audits of European control laboratories. He is also member of the Ph. Eur. Group 15 on Vaccines and Sera and recently joined the mRNAVAC working party at the EDQM. He is also in firm collaborations with the World Health Organization (WHO) as temporary advisor in the framework of the regulatory strengthening of regulatory authorities and Product Summary File evaluation.

Dr Van Molle holds a Master in Biotechnology and obtained a PhD in Molecular Biology from the University of Ghent, Belgium and worked for over 10 year in the academic field, followed by a 2 year period at the Belgian regulatory authority before joining Sciensano.