International Alliance for
Biological Standardization

Globally Harmonized Specifications: Current State and Future Opportunities
January 10-12, 2023
Kunstmuseum, Basel, Switzerland

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Koen Brusselmans received a master degree in Bioengineering Sciences from the University of Leuven (Belgium) in 1996. Afterwards he started a PhD in medical sciences in the laboratory of Transgene Technology of Prof. Dr. Peter Carmeliet and Prof. Dr. Désiré Collen (University of Leuven), which focused on the role of hypoxia-inducible factors in the induction of angiogenesis during embryonic development and tumorigenesis.

After having obtained his PhD in 2001, he worked for 7 years as a post-doctoral fellow in the laboratory of Prof. Dr. Johan Swinnen and Prof. Dr. Guido Verhoeven (University of Leuven) on a research project studying the role of lipogenesis in cancer.

In 2008, he joined the group of 'Quality of vaccines and blood products' at Sciensano (the former Institute of Public Health, Brussels, Belgium), where he is currently working as senior quality assessor for biological medicines. He is involved in assessment of scientific advices and registration files of biological medicines (including vaccines, plasma-derived products and recombinant proteins), in collaboration with the Belgian Medicines Agency (FAMHP) and the European Medicines Agency (EMA). He also participates as expert in GMP inspections of manufacturers of biotech products and plasma-derived products.

Koen Brusselmans is Plasma Master File coordinator and alternate member for Belgium in the CHMP Biologics Working Party at the EMA.