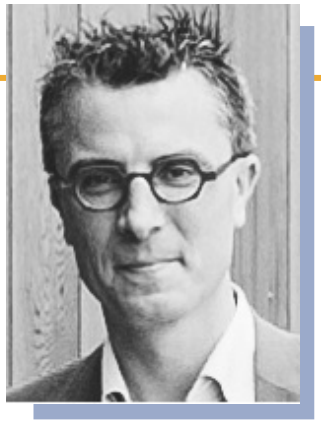
International Alliance for Biological Standardization

BIOSKETCH

Autogenous Vaccines Hybrid Meeting

Quality of Production and Movement In a Common Market

September 14 – 16, 2021



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Dries qualified as a pharmacist and holds an additional degree of Master in Business Planning. He started working in regulatory affairs for human medicinal products in 2003 in the pharmaceutical company Pfizer. In 2004 he made the shift to the Belgian Federal Agency on Medicines where he started working as project manager for veterinary medicinal products. Since 2010 Dries is the Head of the veterinary division within the Federal Agency on Medicines and Health Products. As delegate in the council working party on veterinary medicines he was also involved in the elaboration of the Regulation 2019/6/EC on veterinary medicines. Dries is also the Belgian delegate in multiple EU committees and working groups such as the Standing committee, pharmaceutical committee, CMDv, QRD and NtA. Since 2019 Dries is also the chair of the CMDv Legislation working group. This working group is responsible to coordinate the implementation of the Regulation at the level of the CMDv