Curriculum vitae (short form):

Jobst Limberg is a scientific director heading the unit "Scientific Quality in European Procedures" in the department for European and International affairs at BfArM. He is the German member of the Quality Working Party of the EMA responsible for scientific coordination of pharmaceutical quality in the German Drug Regulatory Agency.

Key issues of his evaluation experience have been analytical and technological topics. Also he participates in the committee pharmaceutical technology of the German Pharmacopoeia and has been involved in harmonisation of several general methods of the European Pharmacopoeia. Additionally he takes an active part in the development of national and international guidelines. Currently he is also involved in the national PAT group and the respective international groups at EMA in London and EDQM in Strasbourg.

After receiving his Ph.D. in pharmaceutical technology he worked as a Post Doc in the analytical laboratories of the University of Florida, USA. He was responsible for the development of methods and routine analysis of drugs and their metabolites in biological materials. Starting in 1990 he joined BfArM as a quality assessor; from 1992 to 1995 he worked as a group leader in the laboratories of the agency; from 1995 to 2005 he was head of the unit "Pharmaceutical Technology". From 2005 to 2011 he has been head of the interdisciplinary regulatory unit "Cardiology".