



International Alliance for
Biological Standardization

Europe



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- Trained as Biologist, with a PhD in Plant Biochemistry (Leiden University, 1993).
- Postdoc positions in Biochemistry, Biotechnology & modelling
- Assessor of Biopharmaceuticals (since 2002); Specialised in vaccines, allergens, biosimilars, immunogenicity, and Cell & Gene therapy products.
- Chair EMA Quality Innovation Group (QIG; since 2023). The QIG supports the translation of innovative approaches to the design, manufacture and quality control of medicines.
- Chair HMA Substance Validation Group (SVG; since 2020). The SVG is involved in development and maintenance of EU-SRS (EU-Substance Registration System); a database with all substances in medicinal products with their molecular structure.
- Involved in various academic research projects to support the assessment of (bio)pharmaceuticals and other regulatory activities.