BUREAU VERITAS
Certification

Certificate

For

KAHLITY
GROW YOUR BUSINESS

KAHLITY GmbH

Bruderholzallee 53, 4059 Basel / Switzerland
with subsidiaries according to appendix

Bureau Veritas Certification hereby confirms that the management system of the above-mentioned organisation has been assessed and complies with the requirements set out in the following standards/regulations.

Standards/Regulations:

SN EN ISO 13485 :2016

The management system comprises:

Swiss and European Authorized Representative, Importer, and Consulting services for the medical device industry, as well as for pharmaceutical companies designing, developing, manufacturing, or ensuring compliance for drug-device combination (DDC) products.

Date of initial certification:	**08.07.2024**	End of the last certification cycle:	--
Date of certification:	**28.05.2024**	Start of the new certification cycle:	**08.07.2024**

The requirements of the standards/regulations must be continuously fulfilled throughout the validity of this certificate. This will be ensured and guaranteed through regular monitoring by Bureau Veritas Certification.
This certificate is valid until: **07.07.2027**

Bureau Veritas Certification will provide information on the validity of this certificate on request at any time. Additional information on the management system and the area of applicability should be obtained from the organisation itself.

Certificate number: **CH19213050** Version: **2** Issue date: **26.05.2025**

SWISS ACCREDITATION
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Appendix

to Certificate N° **CH19213050**, Version **2**

KAHLITY GmbH

Bruderholzallee 53, 4059 Basel / Switzerland

Bureau Veritas Certification has issued this annex to the Management Certificate of the above mentioned company.

Standards/Regulations:

SN EN ISO 13485 :2016

Locations	Field of activity
KAHLITY GmbH Bruderholzallee 53, 4059 Basel / Switzerland	Swiss Authorized Representative, Importer, and Consulting services for the medical device industry, as well as for pharmaceutical companies designing, developing, manufacturing, or ensuring compliance for drug-device combination (DDC) products
KAHLITY France SAS 51 Quai Joseph Gillet, 69004 Lyon / France	European Authorized Representative, Importer, and Consulting services for the medical device industry, as well as for pharmaceutical companies designing, developing, manufacturing, or ensuring compliance for drug-device combination (DDC) products

Issue date: **26.05.2025**