144987-20-05-19

EC DESIGN-EXAMINATION CERTIFICATE

Directive 93/42/EEC on Medical devices, Annex II (4)

CE Certiso Ltd. (NB 2409) certifies that the design of the device concerning to the listed devices and device categories conforms to the requirements of the directive.

Name of the manufacturer:

BioScience GmbH

Headquarters: **Walsmühler Straße 18, D-19073 Dümmer, Germany**

Scope:

Cross-linked hyaluronic acid with dextranomer dermal filler

The certificate covers the following devices:

Description of the device	Type	Intended use	Model	Risk class
Cross-linked hyaluronic acid with dextranomer	Hylan Gel Dermal Filler DX	soft tissue augmentation	CRM DX, Genefill DX	III

This certificate is valid only with the system certificate No. **144983-20-05-19**, in case of successfully conducted annual surveillance audits.

ID number of the related design examination report: **118-G4-190327**

Issue: 1
Issued: 19 May 2020
First issued: 19 May 2020
Start date of certified status: 13 January 2016

Expires:
25 May 2024

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