

**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 Dummer, 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

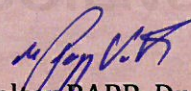
Issued: 19 May 2020

First issued: 19 May 2020

Start date of certified status: 13 January 2016

Expires:

**25 May 2024**

  
Valter PAPP, Dr.  
General Manager

