EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

Naturelize GmbH

Kasselerstraße 47
34308 Bad Emstal
Germany

that the design of the following device(s)

Hyaluronic acid fillers
Inline, Inline Soft, Maxface, Maxface Soft, Pearl, Restore

is conform to the Essential Requirements of Annex I of the Council Directive 93/42/EEC concerning medical devices.
This EC Design Examination Certificate is only valid in connection with the valid DQS Medizinprodukte GmbH Certificate No. 523027 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination: CED-001 File for products „Inline Inline Soft, Maxface, Maxface Soft, Pearl, Restore“ dated 2016-08-16

Further basis for the examination is referenced in the examination report and relating documents mentioned below.

Examination report: Naturlize_Bericht_EGA.doc dated 2016-08-16

The results of the examination are contained in the above mentioned report and the relating documents mentioned within.

Certificate registration no.	523027 MRA
Certificate unique ID	170662074
Effective date	2016-10-13
Expiry date	2021-09-21
Frankfurt am Main	2016-10-13

DQS Medizinprodukte GmbH

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DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.