EC Design Examination Certificate

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

SIC invent AG

Birmannsgasse 3
4055 Basel
Schwitzerland

that the design of the following device(s)

SIC Nature Graft, absorbable inorganic bone substitute material made out of biological hydroxylapatite/ flourohydroxylapatite intended for the augmentation of bony periodontal defects, sinus elevation grafting, treatment of intrabony periodontal defects, and extraction socket grafting
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. 293238 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination:	PHA02_A00_Technische_Doku_Nature_Graft_v1.9 dated 08.12.2017 Further basis for the examination is referenced in the examination report and relating documents mentioned below.
Examination report:	18_SIC_Nature_Graft_EGA_Bericht dated 07.04.2018 The results of the examination are contained in the above mentioned report and the relating documents mentioned within.

Certificate registration no.	293238 MRA
Certificate unique ID	170711502
Effective date	2018-04-15
Expiry date	2023-04-14
Frankfurt am Main	2018-04-15

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Dr. Thomas Feldmann
Head of Certification Body

August-Schanz-Straße 21, 60433 Frankfurt am Main,
Tel. +49 (0) 69 95427-300, medical.devices@dqs-med.de

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

411.23 Version 1.0

Annex to certificate
Certificate registration No.: 293238 MRA
Certificate unique ID: 170711502
Effective date: 2018-04-15

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REF	Description
510808	SIC Nature Graft 0,5 ml
510816	SIC Nature Graft 1,0 ml
510824	SIC Nature Graft 2,0 ml