



## **EC Design Examination Certificate**

Council Directive 93/42/EEC Annex II Section 4

This is to certify that the manufacturer

## **Degradable Solutions AG**

Wagistrasse 23 8952 Schlieren Switzerland

that the design of the following device(s)

easy-graft™ Classic

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. 081238 MR2. Changes to the approved design are subject to further approval by the Notified Body.

Basis of examination:	easy-graft CLASSIC dated September, 2015
	Further basis for the examination is referenced in the examination report and relating documents mentioned below.
Examination report:	Report TFR easy graft classic V01 dated 2016-01-05
	The results of the examination are contained in the above mentioned report and the relating documents mentioned within.

Certificate registration no.	341505 MRA
Certificate unique ID	170639346
Effective date	2016-01-05
Expiry date	2021-01-04
Frankfurt am Main	2016-01-05

## **DQS Medizinprodukte GmbH**

August-Schanz-Straße 21, 60433 Frankfurt am Main,

Tel. +49 (0) 69 95427-300, medical.devices@dqs-med.de

Frank Graichen Managing Director

Dr. Thomas Feldmann Head of Certification Body

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