



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



Product Service

EC Certificate

EC Design-Examination Certificate
Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)
(Devices in Class III)

No. G7 004200 0006 Rev. 00

Manufacturer: **Biocomposites Ltd**
Keele Science Park
Keele, Staffordshire ST5 5NL
UNITED KINGDOM

Product: **Bone substitutes**
Calcium matrix for bone and soft tissue
implantation

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with MDD Annex II (4). The design of the devices conforms to the requirements of this Directive. For marketing of these devices an additional Annex II certificate is mandatory. See also notes overleaf.

Report no.: 713131301

Valid from: 2020-04-14

Valid until: 2024-05-26

Date, 2020-04-14

Christoph Dicks
Head of Certification/Notified Body



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



Product Service

EC Certificate

EC Design-Examination Certificate
Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)
(Devices in Class III)

No. G7 004200 0006 Rev. 00

Model(s): Stimulan® Kit
Stimulan® Rapid Cure

Product Code (Ref)	Description	Size
600-005	Stimulan Kit	5cc
600-010	Stimulan Kit	10cc
620-005	Stimulan Rapid Cure	5cc
620-010	Stimulan Rapid Cure	10cc
620-020	Stimulan Rapid Cure	20cc