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ZLG-BS-244.10.08

TÜV SÜD 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 104609 0002 Rev. 02

Manufacturer:

CERUS Corporation
1220 Concord Avenue
Concord CA 94520
USA

Product:

Blood Processing Devices
Pathogen Inactivation Disposables

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.: 713164217

Valid from: 2020-03-13
Valid until: 2024-05-26

Date, 2020-03-13

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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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 104609 0002 Rev. 02

Model(s): **INTERCEPT Blood System for Platelets (Amotosalen Photochemical Treatment)**

Parameter:

INTERCEPT Processing Set for Small Volume Platelet Units (INT21)

INTERCEPT Processing Set for Large Volume Platelet Units (INT22)

INTERCEPT Platelet Processing Set with Dual Storage Containers (INT25)

INTERCEPT Platelet Processing Set with Triple Storage Containers (INT26)

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