



Product Service

EC Design Examination Certificate

(Annex II, section 4 of the Directive 93/42/EEC on Medical Devices)

No. G7 10 05 74018 003

Manufacturer: **CERUS Corporation**
2550 Stanwell Drive
Concord CA 94520
USA

EC-Representative: **Cerus Europe B.V.**
Stationsstraat 79-D
3811 MH Amersfoort
THE NETHERLANDS

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 aforementioned devices according to Annex II, section 4 of the Directive 93/42/EEC on Medical Devices. The design of the devices conforms to the provisions of this Directive. For marketing of these products an additional Annex II.3 certificate is mandatory. See also notes overleaf.

Report no.: 71369581

Valid until: 2012-05-29

Date, 2010-06-22

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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

Parameters:

- INTERCEPT Processing Set for Apheresis Platelets
- INTERCEPT Processing Set for Pooled Buffy Coat Platelets
- INTERCEPT Processing Set for Small Volume
- INTERCEPT Processing Set for Large Volume
- INTERCEPT Platelet Processing Set with Dual Storage Containers

Facility(ies):

Cerus Europe B.V.
Stationsstraat 79-D, 3811 MH Amersfoort, THE NETHERLANDS

CERUS Corporation
2550 Stanwell Drive, Concord CA 94520, USA