



APPROVAL
EC Directive 93/42/EEC Annex II, Article 3
Full Quality Assurance System
Medical Devices

Registration No.: HD 60013655 0001

Report No.: 30491925 001

Manufacturer: Thermogenesis Corporation
2711 Citrus Road
Rancho Cordova, CA 95742
USA

Scope: Design, Development, Manufacturing and Servicing of
Medical Devices

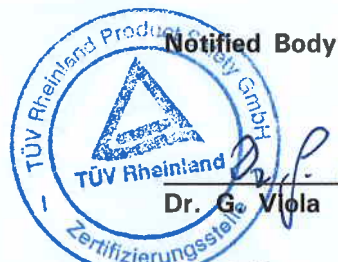
Products: see attachment

Replaces Approval, Registration No.: HD 60009010 0001

Date of Expiry: 21.03.2011

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex II, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex II, Article 5 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Cologne, 22.03.2006



TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. CE

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to
Registration No.: HD 60013655 0001
Report No.: 30491925 008

Manufacturer: ThermoGenesis Corporation
2711 Citrus Road
Rancho Cordova CA 95742
USA

Scope:

Products:

- Freezers for Biological Material
BioArchive, AXP System
- Freezers for Blood Plasma
- Fibrin Sealant System
CryoSeal FS System
TPD
- MarrowXpress System
- Res-Q60 Disposable Kit
- Art 21 Disposable Kit

Date 16.03.2010

