

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 614420
Issued To: **ThermoGenesis Corp.**
2711 Citrus Road
Ranch Cordova
California
95742
USA

In respect of:

Design, development and manufacture of blood and bone marrow processing systems and fibrin processing systems.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):



Stewart Brain, Head of Compliance & Risk -
Medical Devices

First Issued: **2015-01-13**

Date: **2017-09-06**

Expiry Date: **2021-01-13**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

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95742
USA

Subcontractor:

Service(s) supplied

Biotest Laboratories, Inc.
9303 West Broadway Ave
Brooklyn Park
Minnesota
55445
USA

**Control of Sterilization
Manufacture**

MDSS GmbH
Schiffgraben 41
Hannover
30175
Germany

EU Representative

Synergy Health AST, SRL
B16, Street 4, Avenue O
El Coyol Free Zone
El Coyol
Alajuela
20102
Costa Rica

E beam Sterilization

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Vention Medical Costa Rica
Parque Zona Franca Metropolitana
Edificio 2C
Barreal de Heredia
40101
Costa Rica

**Control of Sterilization
Manufacture**

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EC Certificate - Full Quality Assurance System Certificate History

Certificate No: **CE 614420**
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Date	Reference Number	Action
13 January 2015	8153014	First issue. Transfer from another Notified Body.
26 October 2015	8351134	Change of Legal manufacturer name as 'ThermoGenesis Corp. (Cesca Therapeutics Inc.)'
11 January 2016	8405704	Renewal. Reduction of scope to 'Design, Development and Manufacture of Freezers for Biological Materials, Stem Cell and Bone Marrow Processing Systems and Fibrin processing systems.' Inclusion of reference to sterile accessories associated with the Bone marrow processing system in the scope. Addition of 'Control of Sterilization' activity to significant subcontractors, Vention Medical. Addition of 'Synergy' and 'Steris' as significant subcontractors for sterilization activities.
30 August 2016	8573919	Change of Legal manufacturer name to 'Cesca Therapeutics Inc. Dba ThermoGenesis Corp'
Current	8676453	Removal of subcontractors: - Vention Medical Inc - Steris Isomedix Services Reduction in scope to remove 'Those Aspects of....' and 'Freezers for Biological Materials' Rewording of scope to replace stem cells with blood. Company name change to ThermoGenesis Corp