EC Type Examination Certificate

for the EC Conformity Assessment according to the Directive 97/23/EC of the European Parliament and of the Council of 29 May 1997 on the Approximation of the Laws of the Member States concerning Pressure Equipment, in its valid version.

Certificate No.: **88 402-13 HH**

Authorized Representative:
Emergo Europe
Molenstraat 15
2513 BH The Hague
The Netherlands

Manufacturer:
Groupe Medical Gaumond Inc.
2500 rang 4
Saint Calixte, Quebec
J0K 1Z0 CANADA

Certification Object

Manufacturers Designation:	HematoCareTM (transportable Hyperbaric Chamber)
Description:	Assembly (including flexible Pressure Vessel)
Category:	IV
Applied Module:	B
Applied Standard:	./.

Basis:
Basis for Examination was the EC Directive 97/23/EC.

Results:
The above mentioned Pressure Equipment is in accordance with the relevant essential safety requirements of the EC Directive 97/23/EC (Examination Report No. 885/13).

Accompanying Documents:
- GL Letter with Ref. No10-018974 dated 2013-06-21 as well as the technical documents of the Pressure Equipment mentioned in this letter.
- GL Letter with Ref. No10-018974 dated 2012-08-30 as well as the technical documents of the Pressure Equipment mentioned in this letter.

EC Type Examination Certificate valid until: 2023-06-20

Hamburg, 2013-06-21

Germanischer Lloyd

M. Schmidt

Matthias Schmidt

Notified Body for the Certification of Pressure Equipment
Identification No. 0098

Germanischer Lloyd SE, Post Box 11 16 06, 20416 Hamburg, Germany, Telephone +49 40 36149-0, Fax +49 40 36149-200, E-mail headoffice@gl-group.com