

Declaration of Conformity MDD, TPED, RoHS 2



Legal Manufacturer / Address:	CAIRE Inc.	
	2200 Airport Industrial Dr, Suite 500	
	Ball Ground, GA 30107 USA	
Manufacturing Facilities	CAIRE Inc.	
2.5	2205 Airport Industrial Drive	
	Ball Ground, GA 30107	
	USA	
Repair Facilities/Final Configuration/Ten	CAIRE Medical Italy Srl Via 10	CAIRE Medical Germany GmbH
Year Retest	Canada	Arnold-Höveler-Strasse 2,
	35127	40764 Langenfeld. Germany
	Padova, ITALY	
	CAIRE Medical Ltd. Unit 6, Ashville way	
	Wokingham RG41 2PL UK	
Notified Body	TPED: Apragaz A.S.B.I. (0029)	MDD: GMED SAS (0459)
	Chaussée de Vilvorde, 156	1, rue Gaston Boissier 75015
	1120 Bruxelles, Belgium	PARIS, France
TPED/MDD Representative	Medical Product Service GmbH (MPS)	
	Borngasse 20	
	35619 Braunfels, Germany	
Product Families	Portable Liquid Oxygen Units (Stroller, Sprint, Spirit, HELiOS)	
	Base Liquid Oxygen Units (Liberator, HELiOS)	
MDD Equipment Classification	Class IIa, MDD Appendix IX, Rule 11	
Global Medical Device Nomenclature	35910-Portable Liquid Oxygen Units	
(GMDN) Code:	35221-Base Liquid Oxygen Units	
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Start of CE Marking	Helios base: 11 November 2008	
	Helios portable 30 November 2001	
Other CD:	Stroller, Sprint, Spirit, Liberator: 13 December 2002	
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	Helios portable 30 November 2001	
	Stroller, Sprint, Spirit, Liberator: 13 December 2002	

This declaration applies to all CE marked devices manufactured from date of issuance until it is either superseded by another declaration or withdrawn. I, the undersigned, hereby declare the equipment specified above and its CAIRE approved critical components meets the essential requirements and conforms to the following, as certified by Apragaz A.S.B.I., NB 0029 and GMED SAS, NB 0459.

- Council Directive 93/42/EEC, Medical Devices Directive and Amendment 2007/47/EC Annex II, Clause 3 (Certificate 31275)
- EN-ISO 13485:2016, Quality Management Systems (Certificate 31276)

These devices also conform to the following Directive(s):

- Council Directive 2011/65/EU, Restriction of hazardous substances
- Closed cryogenics oxygen receptable made according to the Transportable Pressure Equipment Directive 2010/35/EU and covered by the following certificates:
 - Surveillance of the Applicants in-house inspections: 1.8.7.3 & 1.8.7.4 of ADR(2019): Certificate 09/US/1952

Edward Kim, Vice President – Engineering and Regulatory

Date

2/4/2021