

ATTESTATION CE / EC CERTIFICATE

Approbation du Système Complet d'assurance Qualité / Approval of full Quality Assurance System

ANNEXE II excluant le point 4 Directive 93/42/CEE relative aux dispositifs médicaux

ANNEX II excluding section 4 Directive 93/42/EEC concerning medical devices

Pour les dispositifs de classe III, un certificat CE de conception est requis

For class III devices, a EC design certificate is required

Fabricant / Manufacturer

CAIRE Inc.

2200 Airport Industrial Drive, Suite 500

BALL GROUND, GA 30107 UNITED STATES

Catégorie du(des) dispositif(s) / Device(s) category

**Systèmes d'oxygène liquide et leurs composants à utiliser avec des produits de santé à domicile
en thérapie respiratoire et concentrateurs d'oxygène**

*Liquid oxygen systems and their components for use in home healthcare products for respiratory therapy
and oxygen concentrators*

Voir détails sur addendum / See attachment for additional information

GMED atteste qu'à l'examen des résultats figurant dans le rapport référencé T000180ER, le système d'assurance qualité - pour la conception, la production et le contrôle final - des dispositifs médicaux énumérés ci-dessus est conforme aux exigences de l'annexe II excluant le point 4 de la Directive 93/42/CEE.

GMED certifies that, on the basis of the results contained in the file referenced T000180ER, the quality system - for design, manufacturing, and final inspection - of medical devices listed here above complies with the requirements of the Directive 93/42/EEC, annex II excluding section 4

La validité du présent certificat est soumise à une vérification périodique ou imprévue

The validity of the certificate is subject to periodic or unexpected verification

Début de validité / Effective date : **April 27th, 2020 (included)**

Valable jusqu'au / Expiry date : **May 26th, 2024 (included)**



On behalf of the President

Béatrice LYS

Technical Director

DocuSigned by:
Béatrice Lys
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Identification des dispositifs / Identification of devices

Description du Dispositif Médical <i>Medical Device Description</i>	Classe du Dispositif Médical <i>Medical Device Class</i>
VisionAire	IIa
Newlife Family	
Eclipse	
Spirit	
Helios Range	
Sprint/Stroller	
Liberator	
Freestyle Comfort	
Flowmeter, SureFlow	
Saros 4000	
Companion 5	

Les produits couverts par ce certificat sont référencés sur la liste des produits par le GMED en date du 5 Mars 2020.

The products covered by this certificate are listed on the product list authenticated by GMED on March 5, 2020

Ce certificat couvre les sites et activités suivants /

This certificate covers the following sites and activities:

CAIRE Inc.

2200 Airport Industrial Drive, Suite 500
 Ball Ground, GA 30107
 USA

Siège social & conception

Headquarters & design



GMED 0459

On behalf of the President

Béatrice LYS

Technical Director

DocuSigned by:
Beatrice Lys
 EF33BDABBA0A3...



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Addendum of the certificate n° 31275 rev. 12
Dossier / File N° T000180ER

CAIRE Inc.

2205 Airport Industrial Drive
Ball Ground, GA 30107
USA

Fabrication & contrôle final

Manufacturing & final control

CAIRE Medical Technology (Chengdu) co., Ltd.

No. 48 Qingma Rd, South Section,
Chengdu Modern Industrial Park
Pidu district, Chengdu, Sichuan 611730
China

Fabrication & contrôle final

Manufacturing & final control

CAIRE Medical Germany GmbH

Arnold-Höveler Strasse 2
40764 Langenfeld
Germany

Fabrication & contrôle final

Manufacturing & final control

4 Sites/ 4 Sites

GMED	0459
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On behalf of the President

Béatrice LYS

Technical Director

DocuSigned by:

Béatrice Lys

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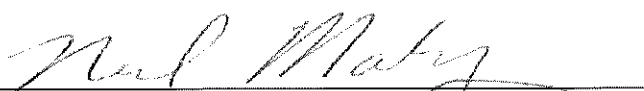
Legal Manufacturer / Address	CAIRE Inc. 2200 Airport Industrial Dr, Suite 500 Ball Ground, GA 30107, USA	
Manufacturing/Repair Facilities	CAIRE Inc. 2205 Airport Industrial Drive Ball Ground, GA 30107 USA	CAIRE Medical Technology (Chengdu) Co., Ltd. No. 48 Qingma Road, South Section Chengdu Modern Industrial Park Pidu district, Chengdu 611730, Sichuan China
Repair Facilities	CAIRE Medical Italy Srl Via 10 Canada 35127 Padova, Italy	CAIRE Medical Ltd. Unit 6- Ashville Way Wokingham, Berkshire RG41 2PL United Kingdom
Repair, distribution & order fulfilment, final configuration	CAIRE Medical Germany GmbH Arnold-Höveler-Strasse 2, 40764 Langenfeld. Germany	
Notified Body	GMED SAS (0459) 1, rue Gaston Boissier 75015 PARIS France	
Authorized Representative	Medical Product Service GmbH (MPS) Borngasse 20 35619 Braunfels, Germany	
Product Families	Oxygen Concentrators (Eclipse Oxygen System, NewLife Intensity 10, NewLife Elite, VisionAire, Saros 4000, Companion 5, and FreeStyle Comfort)	
MDD Device Classification	Class IIa (MDD Appendix IX, Rule 11)	
Global Medical Device Nomenclature (GMDN) Code	12873- Stationary Oxygen Concentrators 31321-Portable Oxygen Concentrator	
Start of CE Marking	Eclipse Oxygen System: 14 March 2007 NewLife Intensity 10: 08 October 2008 NewLife Elite: 22 April 2019 VisionAire: 02 June 2008 FreeStyle Comfort: 01 March 2018 Saros 4000: 05 March 2020 Companion 5: 05 March 2020	

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 meets the essential requirements and conforms to the following, as certified by GMED SAS, NB 0459.

- Council Directive 93/42/EEC, Medical Devices Directive (Annex II, excluding section 4) and Amendment 2007/47/EC (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


Neal Maloy, Director – Quality and Regulatory

April 24, 2020

Date

Identification des dispositifs / Identification of devices

Description du Dispositif Médical <i>Medical Device Description</i>	Classe du Dispositif Médical <i>Medical Device Class</i>
FreeStyle	Ila
Focus	Ila
VisionAire	Ila
Newlife Family	Ila
CR50 O2 Regulator	Ilb
Companion 5	Ila
eQuinox	Ila
FreeStyle	Ila
Eclipse	Ila
SAROS	Ila
Sureflow	Ila
Spirit	Ila
Companion Range	Ila
Helios Range	Ila
Sprint/Stroller	Ila
Liberator	Ila

Les produits couverts par ce certificat sont référencés sur la liste des produits authentifiée par le LNE/G-MED en date du 18 août 2016.

The products covered by this certificate are referenced on the list of products authenticated by LNE/G-MED on August 18, 2016.



LNE/G-MED

0459

**On behalf of the Certification Director
Pascal PRUDHON
Multifields Certification Division Manager**

ADD

720 DM 0701-31 rev 5 du 28/07/2015

Ce certificat couvre les sites suivants / This certificate covers the following sites:

CAIRE Inc.
2200 Airport Industrial Drive, Suite 500
Ball Ground, GA 30107
USA

Siège social & conception
Headquarters & design

CAIRE Inc.
2205 Airport Industrial Drive
Ball Ground, GA 30107
USA

Fabrication & contrôle final de produits de santé à domicile
Manufacturing & final control of home healthcare products

CAIRE Inc.
500 Commerce Drive
Buffalo, NY 14228
USA

Conception, fabrication & contrôle final de concentrateurs d'oxygène
Design, manufacturing & final control of oxygen concentrators

CAIRE Inc.
12230 World Trade Drive Suite 100
San Diego, CA 92128
USA

Conception, fabrication & contrôle final des produits de santé à domicile
Design, manufacturing & final control of home healthcare products



LNE/G-MED

0459

On behalf of the Certification Director
Pascal PRUDHON
Multifields Certification Division Manager

ADD

720 DM 0701-31 rev 5 du 28/07/2015



Le progrès, une passion à partager

Certification
Médical-Santé

Addendum au certificat n° 31275 rev. 2
Addendum of the certificate n° 31275 rev. 2
Dossier / File N° T000103 -T & T000103-3-DOCR

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Chart Biomedical (Chengdu) Co. Ltd.
No. 48 Qingma Rd, South Section,
Chengdu Modern Industrial Park
Pixian, Chengdu 611730, Sichuan
China

Fabrication & contrôle final de produits de santé à domicile
Manufacturing & final control of home healthcare products

Chart Biomedical GmbH
Essener Strasse 68
42327 Wuppertal
Germany

Fabrication & contrôle final de produits de santé à domicile
Manufacturing & final control of home healthcare products

6 sites / 6 sites

LNE/G-MED	0459
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On behalf of the Certification Director
Pascal PRUDHON
Multifields Certification Division Manager

ADD

720 DM 0701-31 rev 5 du 28/07/2015

Laboratoire national de métrologie et d'essais • établissement public à caractère industriel et commercial
LNE/G-MED • Organisme notifié n° 0459
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