EC DECLARATION OF CONFORMITY

We, **Air Liquide Medical Systems S.A.**
Parc de haute technologie
6, rue Georges Besse
92182 ANTONY Cedex
FRANCE

declare under our sole responsibility that the product range:

XO

Valve Integrated Pressure Regulator (VIPR), and its accessories, comply with the requirements of the European Directive 93/42/EEC concerning medical devices, with all amendments and transposing legislation, and all others Union legislation applicable to the device.

Notified Body: GMED (0459)
1, rue Gaston Boissier - 75724 PARIS Cedex 15 – France

Annex II conformity certificate, references n° GMED-33855 (DFM).

Classification: class IIb, according to rule 11 as specified in annex IX of the Directive 93/42/EEC.

Designation	Reference	Date of first issue
XO O2 200B 25E OL DIN 15L	CK062539	14/03/2017
XO O2 200B 25E OL NF 15L	CK062538	14/03/2017
XO O2 200B 25E OL UNI 15L	CK062540	01/04/2020
XO O2 200B 25E OL SS 15L	CK062541	01/04/2020
XO O2 200B 25E OL CARBA 15L	CK062542	01/04/2020
XO O2 200B 25E OL SIS 15L	CK062543	23/05/2023
XO O2 200B 25E OL DISS 15L	CK062544	26/10/2022
XO SMART O2 200B OL NF 15L	CK062548	01/04/2020
XO SMART O2 200B OL DIN 15L	CK062549	01/04/2020
XO SMART O2 200B OL CARBA 15L	CK062552	01/04/2020
XO SMART O2 200B OL SS 15L	CK062551	01/04/2020
XO SMART O2 200B OL UNI 15L	CK062550	01/04/2020
XO SMART O2 200B OL SIS 15L x 10	CK062553	23/05/2023
XO SMART O2 200B OL BS 15L x 10	CK062554	23/05/2023
XO SMART O2 200B OL DISS 15L x 10	CK062556	26/10/2022

This EC Declaration of Conformity is valid for the placing on the market or the putting into service in the countries within the European Union and the European Economic Area where the following languages are authorized by the national competent authority:

FR	Français	**EN**	English	**DA**	Dansk	**DE**	Deutsch
ES	Español	**IT**	Italiano	**NO**	Norsk	**PL**	Polski
PT	Português	**SV**	Svenska	**NL**	Nederlands		

At Antony, on 26-oct.-2023

Olivier VAROMME
Quality and Regulatory Affairs Director

DocuSigned by:
Olivier VAROMME
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