EU DECLARATION OF CONFORMITY (DoC)

Manufacturer: AHA Hyperbarics GmbH
Registered place of business: Mellergasse 4, 1230 Vienna, Austria
e-mail: ask@aha-hyperbarics.com

SRN: AT-MF-000023416

We hereby declare that the DoC is issued under our sole responsibility and applies to the following devices and products:

Classification: Class IIb
EMDN code / Device Group: Z120308 - **HYPERBARIC CHAMBERS**
Models: AHA Flex 16 V2, AHA Flex 18 V2, AHA Flex 20 V2 and AHA Flex 30
Intended Purpose: AHA Flex hyperbaric chambers are intended to perform hyperbaric oxygen therapy.

Classification: Class IIb
EMDN code / Device Group: Z12159004 - **OXYGEN CONCENTRATORS**
Models: AHA Oxy 22 and AHA Oxy 30
Intended Purpose: AHA Oxy oxygen concentrators are intended to perform hyperbaric oxygen therapy (to be used with the hyperbaric chamber).

UMDNS: 12061

Main Regulation:
Regulation (EU) 2017/745 on Medical Devices (according to Annex IX., ex. 5). As follows the stated products are declared as **medical devices.**

Standards:
The company has established general quality management standard: EN ISO 13485:2016

Notified Body:	TÜV SÜD Product Service GmbH Ridlerstraße 65, 80339 München, Germany
CE mark:	CE 0123
Certificate Nr.:	G15 104291 0006 Rev. 00
Certificate issued:	2026-07-23
Certificate valid:	2031-07-22

Vienna, 23.07.2026
Valid until: 22.07.2031

AHA Hyperbarics GmbH
General Manager
Peter Kokalj