XENIOS | A FRESENIUS MEDICAL CARE COMPANY

EU Declaration of Conformity

according to the Regulation (EU) 2017/745 on Medical Devices (MDR), Annex IV

We, the manufacturer

Xenios AG
Im Zukunftspark
74076 Heilbronn
Deutschland

SRN: DE-MF-000006233

declare herewith under our sole responsibility

that the medical devices

MultiSupport Ground

Basic UDI DI: 4057224-0000-0000-0MSG-X5

falling into the category of

MSG Holder	**Ref. No. 30000285**
MSG Base Plate	**Ref. No. 30000286**
MSG Crossbar	**Ref. No. 38350319**
MSG Backup Module	**Ref. No. 38350318**

further accessories :

MSG Bracket	**Ref. No. 38350320**
MSG Pin 2400	**Ref. No. 38350322**

With the Intended Purpose:

The MSG is intended for intra- and inter-hospital transport of intensive care patients who require extracorporeal gas exchange or extracorporeal circulatory support.
The MSG is only approved for ground transport and accepts the patient kit components (oxygenator, DP3 pump head) as well as the technical components (pump drive, sensor box and console) during transport.
The MSG is intended for use in a hospital environment and for transport in emergency services.

of risk class

I (according to rule 1 set out in annex VIII of Regulation (EU) 2017/745)

CE-marked with

CE

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to which this declaration relates, meet(s) all the provisions of Regulation (EU) 2017/745 of the European Parliament and of the Council on Medical Devices which apply to it.

Commons Specification IEC 60601-1-12, IEC 60068-2-27, IEC 60068-2-64

This EU declaration of conformity is valid latest until **24. August 2026.**

Heilbronn, 24. August 2021

ppa. Christian Peis
Person Responsible for Regulatory Compliance
Xenios AG

ppa. Dr. Knut Jöchle
VP Product Life Cycle
Xenios AG