

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

further accessories :

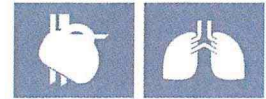
MSG Bracket **Ref. No. 38350320**
MSG Pin 2400 **Ref. No. 38350322**

With the The MSG is intended for intra- and inter-hospital
Intended Purpose: 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

