



**EU DECLARATION OF CONFORMITY**

Annex IX, chapter I and III of Regulation 2017/745 according to  
EU certificate No. G11 037258 0015 Rev. 01 and Expiration Date 2027-07-31  
issued by Notified Body TÜV SÜD Product Service GmbH,  
Ridlerstrasse 65, 80339 Munich, Germany, Identification Number 0123

**Transfer Devices**

Refer to attachment  
(Product name)

Refer to attachment  
(Article Number)

Device classification according to Annex VIII, clause 4.2, rule 2

Class  Is  Im  Ir

We

**Fresenius Kabi AG, Else-Kröner-Str. 1  
61352 Bad Homburg, Germany**

Single Registration Number (SRN): DE-MF-000009273

manufacturer of the above products, hereby declare under our sole responsibility for this Declaration of Conformity that the referenced products comply with all relevant provisions of the Medical Device Regulation 2017/745 and any other relevant Union legislation that provides for the issuing of this EU Declaration of Conformity. The products comply with the general safety and performance requirements of Annex I, further applicable standards and/or other normative documents as listed in the applicable technical documentation. All supporting documentation is kept under the premises of the manufacturer.

Bad Hersfeld, 11.03.2026  
Place and date of issue/

  
Dr. Sandra Leipold, Director Regulatory Affairs  
Name (printed letters), position and signature of authorized person/

Valid starting with the original date of the document until product change



**Attachment to EU DECLARATION OF CONFORMITY**

**Transfer Devices**

Basic UDI-DI: 42502737NIV50611s000000BC

Article Number	Product Name	Intended Purpose
2867001	Freka Lipoflow	For withdrawal and admixture of IV solutions (e.g. drugs or lipids)
2870103	Freeflex+ Transfer Adapter	Reconstitute and/or admix drugs prior to administration to the patient. The device may be used for pediatric and adult populations.