
EU Declaration of Conformity with article list



EU DECLARATION OF CONFORMITY

Annex IX, chapter I and III of Regulation 2017/745 according to EC-certificate
No. G10 037258 0012 Rev. 01 and Expiration Date March 09th, 2026
issued by Notified Body TÜV Süd Product Service GmbH,
Ridlerstrasse 65, 80339 Munich, Germany, Identification Number 0123

Product Family Description
Intravenous pole multi-outlet holder

Product name
Refer to attachment

Article number
Refer to attachment


From serial number / Batch number
Refer to attachment

Device classification according to Annex VIII
LINK+ AGILIA range: Rule 12, Class IIb

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.

Place and date of issue/ Brezins, on <i>Feb. 14, 2024</i>	Name (printed letters), position and signature of authorized person/ Laurent KELLER Regulatory Affairs Manager 
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Valid starting with the original date of the document until product change

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Attachment to EU DECLARATION OF CONFORMITY

LINK+ AGILIA range

LINK+ AGILIA

Article number	Product name	Serial number / Batch number	Basic UDI-DI
Z074601	LINK 4+ AGILIA	25773606	4052682AND31012bCF0000WH
Z076601	LINK 6+ AGILIA	25773636	
Z078601	LINK 8+ AGILIA	25773637	