



**FRESENIUS  
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## Self DECLARATION OF CONFORMITY

Particular procedure for systems and procedure packs  
(Article 12, MDD 93/42/EEC)

Enteral Feeding Tubes percutaneous (Freka® PEG Hybrid Set, FR 15)

Class of procedure pack: IIb

Article code: M90800500 (Freka® PEG Hybrid Set, II FR 15)

Medical Devices / products used within the procedure pack

Article Code	Designation	DoC Designation	Class
7601363	Gastropexy Device II	long-term intragastric feeding gastric decompression	IIb
7755643	Freka PEG CH/FR15, ENFit		

We  
Fresenius Kabi AG  
61346 Bad Homburg, Germany

manufacturer of the procedure pack (article code 7601365), hereby declare under our sole responsibility for this Self Declaration of Conformity:

- that the referenced products comply with all relevant provisions of Directive 93/42/EEC, as amended by 2007/47/EC, and its transposition into national laws.
- All the products comply with the essential 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 legal manufacturer.

We as manufacturer of the procedure pack declare that

- (a) the mutual compatibility of the devices is verified in accordance with the instructions and is carried out his operations in accordance with the instructions; and
- (b) We packaged the system or procedure pack and supplied relevant information to users incorporating relevant instructions from the manufacturers; and
- (c) the whole activity is subjected to appropriate methods of internal control and inspection.



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On behalf of Fresenius Kabi AG,

*01.12.2023*

*S. Leipold*

Dr Sandra Leipold  
Senior Manager Regulatory Affairs  
Fresenius Kabi MedTech – BU INS