EC Declaration of Conformity

Name	Matter	Device Destination
Piégeur (FF1022)	Polypropylene	Single-use Trapper intended to block a volume of air to allow appropriate measurement by the chemical sensor

Medical devices conform to the following standards:

NF EN ISO 13485 :2016 : Medical Device – Quality Management System
NF ISO 2859-1 :2000 : Sampling procedures for inspection by attributes — Part 1: Sampling schemes indexed by acceptance quality limit (AQL) for lot-by-lot inspection
NF EN ISO 14971:2019: Medical devices - Application of risk management to medical devices
NF EN 1041 + A1: 2013: Information supplied by the manufacturer of medical devices
EN ISO 15223-1:2016: Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General

I the undersigned, Dounia Benbachir, Quality Manager of the FIM MEDICAL company located at 51 rue Antoine Primat Villeurbanne - FRANCE assure and declare that the medical devices listed above belong to class I (Rule 1) and satisfy the provisions of annex I (Essential Requirements), annex VII (Evaluation of Tabataba® V2 FF1006DTR100 technical file) of the 93/42/EEC directive and its local adaptation (Book II of the public health code).

Villeurbanne, 12/07/2021,
D.BENBACHIR
Quality Manager

CE