

F.L. MEDICAL s.r.l. Unipersonale Via Enrico Mattei, 20 – 35038 TORREGLIA (Padova) – Italy Tel. +39 049 5211475 / +39 049 5212835 - Fax +39 049 5212566 e-mail: <u>info@flmedical.com</u> - web site: <u>www.flmedical.com</u> C.F. e P.IVA 01134840287 - Cap Soc. 90.000 € <u>i.y.</u> Reg. Imp. di Padova n. 21695 - R.E.A. di Padova n. 187254

EU DECLARATION OF CONFORMITY

MANUFACTURER'S NAME	F.L. MEDICAL s.r.l. Unipersonale	
MANUFACTURER'S REGISTERED PLACE OF BUSINESS AND ADDRESS	Via Enrico Mattei, 20 – 35038 Torreglia (PD) - Italy	
MANUFACTURER'S SINGLE REGISTRATION NUMBER (SRN)	IT-MF-000013918	
DEVICE NAME / TRADE NAME	Test tubes for biological liquids collection	
DEVICE CODES	ref.: Annex I to the present Declaration of Conformity	
RISK CLASS AND CLASSIFICATION RULE	Class A non-sterile Rule 5, according to Annex VIII of the Regulation 2017/746.	
INTENDED USE	Test tubes intended for biological liquids collection	
COMMON SPECIFICATIONS	not applicable	
BASIC UDI-DI	8052109520003UB	
NAME, ADDRESS AND IDENTIFICATION NUMBER OF THE NOTIFIED BODY	not applicable	
CERTIFICATE NUMBER	not applicable	
CONFORMITY ASSESSMENT PROCEDURE	Preparation of the technical documentation (ref. Annexes II and III of Regulation 2017/746) and issue of the EU Declaration of Conformity.	
ADDITIONAL INFORMATION	not applicable	
WE DECLARE UNDER OUR OWN RESPONSIBILITY THAT THE DEVICES ABOVE MENTIONED HAVE BEEN PRODUCED IN COMPLIANCE WITH PRODUCT SPECIFICATIONS, OPERATING INSTRUCTIONS AND LABELLING REQUIREMENTS AND THEREFORE MEET THE PROVISIONS OF THE LAWS IN FORCE ON IN VITRO DIAGNOSTIC MEDICAL DEVICES APPLIED FOR THE CONFORMITY ASSESSMENT PROCEDURE. ALL THE SUPPORTING DOCUMENTATION IS RETAINED AT THE ARCHIVES OF MANUFACTURER'S QUALITY MANAGEMENT SYSTEM, UNDER THE RESPONSIBILITY OF RAQ. THIS DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE RESPONSIBILITY OF THE MANUFACTURER.		
PLACE OF DOCUMENTATION STORAGE	Via Enrico Mattei, 20 – 35038 Torreglia (PD) - Italy	
PLACE AND DATE OF ISSUE OF THE PRESENT DECLARATION	Via Enrico Mattei, 20 – 35038 Torreglia (PD) - Italy	
	Date: 25/08/2022	
NAME, JOB TITLE AND SIGNATURE	Date: 25/08/2022 Alessandro Fiore Quality Assurance Manager (RAQ) Signature:	



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ANNEX I – LIST OF CODES

DEVICE CODE / CATALOGUE NUMBER	DEVICE NAME
22634	SEPARMED ® TECH TEST TUBE 13x75 mm IN POLYPROPYLENE WITH SEPARATING
22034	GRANULES WHITE STOPPER AND LABEL
	SEPARMED ® TECH TEST TUBE 12x86 mm IN POLYPROPYLENE WITH SEPARATING
22626	GRANULES WHITE STOPPER AND LABEL
22020	
22/24	SEPARMED ® TECH TEST TUBE 13x75 mm IN POLYPROPYLENE WITH SEPARATING
22634	GRANULES WHITE STOPPER AND LABEL
00/0/	SEPARMED ® TECH TEST TUBE 12x86 mm IN POLYPROPYLENE WITH SEPARATING
22626	GRANULES WHITE STOPPER AND LABEL
22602	SEPARMED ® TECH TEST TUBE 16x100 mm IN POLYPROPYLENE WITH
	SEPARATING GRANULES WHITE STOPPER AND LABEL
22626	SEPARMED ® TECH TEST TUBE 12x86 mm IN POLYPROPYLENE WITH SEPARATING
	GRANULES WHITE STOPPER AND LABEL
21027	PROMED ® TEST TUBE 16x100 mm 10 ml CONICAL, WITHOUT RIM, GRADUATED,
21027	IN POLYPROPYLENE
	PROMED ® TEST TUBE 16x100 mm 10 ml CONICAL, WITH RIM, GRADUATED, IN
21017	POLYPROPYLENE
22744	SEPARMED ® TECH TEST TUBE 13x75 mm IN POLYPROPYLENE WITH SEPARATING
	GEL + COAGULATION ACCELERATOR RED STOPPER AND LABEL
	SEPARMED ® TECH TEST TUBE 12x86 mm IN POLYPROPYLENE WITH SEPARATING
22734	GEL + COAGULATION ACCELERATOR RED STOPPER AND LABEL
32107	PROMED ® TEST TUBE 12x56 mm WITH K3 EDTA x 2,5 ml OF BLOOD VIOLET
	STOPPER
	PROMED ® TEST TUBE 13x75 mm WITH K3 EDTA x 2,5 ml OF BLOOD VIOLET
32109	STOPPER
32116	PROMED ® TEST TUBE 13x75 mm WITH K3 EDTA x 4 ml OF BLOOD VIOLET
	STOPPER
22213	PROMED ® TEST TUBE 12x56 mm WITH 0,25 ml OF SODIUM CITRATE (3,8 %)
	PINK STOPPER FOR 1 ml OF BLOOD (ESR) AND 2,25 ml OF BLOOD (Coagulation)