


EC 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	<i>Sterile test tubes for blood collection</i>
DEVICE CODES	ref.: Annex I to the present Declaration of Conformity
RISK CLASS AND CLASSIFICATION RULE	Other type of IVD IVD not included in Annex II of Directive 98/79/EC, nor self-testing IVD
INTENDED USE	<i>Test tubes for blood collection, for diagnostic test</i>
COMMON SPECIFICATIONS	<i>not applicable</i>
BASIC UDI-DI	<i>not applicable</i>
NAME, ADDRESS AND IDENTIFICATION NUMBER OF THE NOTIFIED BODY	<i>not applicable</i>
CERTIFICATE NUMBER	<i>not applicable</i>
CONFORMITY ASSESSMENT PROCEDURE	Preparation of the technical documentation (ref. Annex III of Directive 98/79/EC) and issue of the EC Declaration of Conformity.
ADDITIONAL INFORMATION	<i>not applicable</i>
<p>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.</p>	
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: 21/04/2023
NAME, JOB TITLE AND SIGNATURE	Alessandro Fiore Quality Assurance Manager (RAQ)  Signature:



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ANNEX I – LIST OF CODES

DEVICE CODE / CATALOGUE NUMBER	DEVICE NAME
50334	"SEDI-RATE" PIPETTE GRADUATED FROM 0 TO 180 mm