

Numero di accettazione:

TEST ITEM DATA SHEET

All.1 MORD en Rev 3 del 28/11/2023

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Study Sponsor (Name and address)			
Name of Test Item			
As it will be reported in the Test Report			
Product Type *			
IUPAC Name (If applicable)			
CAS Number (If applicable)			
For medical device only			
Nature of body Contact	Surface MD	Ext communicating MD	Implant medical device
Rif Table A.1 ISO 10993-1	☐ Intact skin	☐ Blood path indirect	☐ Tissue / Bone
If applicable	☐ Mucosal Membrane	☐ Tissue / bone / dentin	□ Blood
	☐ Breached surface	☐ Circulating blood	
Contact Duration	□ ≤ 24 hours	□ > 24 h to 30 days	□ > 30 days
Surface area and thickness of the device or We	ight of the device		
Physical appearance □ Device □ Solid □ Liquid □ Gel □ Powder			
Code No.			
Lot No.			
Sterility			
Method of sterilization:Sterilization Lot: Batch produced by (Name and address)			
if different from sponsor			
Date of Manufacture			
Date of Expiry / Valid up to			
Date of Retest			
Number of test items			
Storage conditions ☐ Room Temperature ☐ Cool and dry (+2 to +8°C) ☐ Frozen (< -10°C) ☐ Others: Controlled transport condition (if applicable):			
Safety Precautions, if any			
Material safety data sheet attached			□ Yes □ No
Certificate of analysis			□ Yes □ No
GLP management for the requested tests			□ Yes □ No
In the case of non-destructive tests, the sample must be returned (shipping costs paid by the Customer)			□ Yes □ No
Any additional information (optional)			
Name of The Sponsor's Representative			
Signature and date			
*e.g. Agrochemical, Pharmaceutical, Biotech, Industrial Chemicals, Cosmetics, Nutraceuticals, Food and Feed additives, Medical devices others (Specify)			

To be filled by Lab4LIFE

Verificato da: