Bio-Optica
Improving Pathology

Data Sheet

CE

CONTAINERS for HISTOCOLD

Containers for preservative solution for a safe transport of histological specimens from the surgery room to the PA lab
(In conformity with the European Regulation 605/2014)

CODE	DESCRIPTION	PACKAGING	UDI-DI
05-100401	Multipurpose containers 125 ml	250 pcs	08034120276887
05-100402	Multipurpose containers 250 ml	200 pcs	08034120276894
05-100403	Multipurpose containers 500 ml	100 pcs	08034120276207
05-100404	Multipurpose containers 1000 ml	100 pcs	08034120276900
05-100405	Multipurpose containers 3000 ml	50 pcs	08034120276214
05-100406	Multipurpose containers 5000 ml	20 pcs	08034120276917

IVD

In vitro Diagnostic – Medical Device
IVD **Class A**, Reg. UE 2017/746

Basic UDI: 080341202W05020104BB

Manufacturer: Bio-Optica Milano S.p.A.

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Disposable

Date of issue: 16/05/2022
Rev. 001

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Containers for transport of histological samples immersed in Histocold preservative solution.

GENERAL FEATURES

Polypropylene shockproof containers with large open and pressure watertight lid.
Provided with label for the sample identification.

Technical data

Technical data	Chemical composition	Polypropylene	
	Dimensions	125 ml	62 x 55 x 51 mm
		250 ml	88 x 78 x 62 mm
		500 ml	108 x 97 x 78 mm
		1000 ml	120 x 100 x 130 mm
		3000 ml	182 x 172 x 137 mm
		5000 ml	208 x 192 x 192 mm
	Lid	Pressure	
	Colour	Transparent	
Packaging	Primary container	Carton box	
	Secondary container	None	
Storage	Storing conditions	Particular conditions are not necessary for this kind of material.	
	Stability	After the first opening, the product is stable.	
	Validity	Not applicable to this kind of product	
Warning and precautions	Instruction of use	Not requested for this product.	
	Classification of the product	The product is intended for professional laboratory use for healthcare professionals. The product is not chemically dangerous. No warning measures and precautions are required.	
	Disposal	Destroy by thermo-destruction.	
	Recommendations	In the event of a serious accident, we recommend that you immediately inform Bio-Optica Milano S.p.A. and the competent authorities.	

REVISION N°	REASON	REVISION DATE
001	Regulation adjustment UE 2017/746 - IVDR	16/05/2022

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