



## CRYO - SPRAY



CODE	DESCRIPTION	VOLUME	PACKAGING
08-SPRAY	Freezing medium spray for histology Cryo - Spray	150 ml	12 spray bottles



In vitro Diagnostic – Medical Device  
IVD Class A, Reg. UE 2017/746  
UDI-DI: 08033976236687  
Basic UDI: 080339762W01030799Y5



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

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Rev. 001

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Freezing medium spray for histology.

Cryo – Spray does not contain fluorinated greenhouse gases, so the Regulation (EU) 842/06 article 7 par 2 is not applicable.

Not flammable.

### GENERAL FEATURES

The product quickly freezes tissues before cutting with cryostat and cools embedded samples before microtome cut, where the technicians have not a cold plate.

Spray directly on the sample's surface.

### Technical data

Specifications	Component	HFO-1234ZE
	Colour	Colourless
	Odour	Characteristic
Packaging	Primary container	Spray bottle
	Label	Serigraphed surface
	Secondary container	Carton box, white colour
Storage	Storage	Store in a place where adequate ventilation is ensured, away from direct sunlight at a temperature below 50°C / 122°F, away from any combustion sources.
	Storage temperature	Store at a temperature below 50°C / 122°F.
	Stability	After the first opening, the product is usable until the expiry date, if correctly stored.
	Validity	5 years from the manufacturing date
Warning and precautions	Product classification	The product must be used by specialized technical operators. Always refer to the safety data sheet for information on the risks of the mixture, the precautionary measures during use, and the measures first aid and in case of accidental release. Do not use if the primary container is damaged.
	Disposal	Observe all state and local environmental regulations regarding waste disposal. Bestow not used Cryo-spray to the authorized and specialized establishments, in according with the legislation about the disposal of spray bottles.
	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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