EPR-7012-B

TECHNICAL SHEET

GENERAL INFORMATION

Category:
Sterile sampling bags

Family: Twirl'Blue

Lifespan: 5 years

TECHNICAL DESCRIPTION

The Twirl'Blue bags are blue in colour and have a practical and easy-to-use closing system. They are made of a flexible and durable plastic.

SPECIFIC INFORMATION

ITEM

ITEM	Bag
Material :	Polyethylene blend
Color :	Transparent
Dimension :	178 x 305 mm / 7 X 12 po
Thickness :	3 mil.in / 0.0762 mm / 76.2 micron
Total volume :	2200 ml / 75 oz
Utility volume :	1300 ml / 44 oz
Printing type :	Writing area
Opening system :	Perforated line
Closing system :	Attachment with 2 round wires
Sterile :	Yes
End of product life :	Recyclable

PACKAGING INFORMATION

Outer box dimension : (W x D x H)	14.13 po x 9.50 po x 16.25 po 36 cm x 24 cm x 41 cm
Box weight:	22.00 LB / 9.98 KG
Conditioning:	1000 (4 x 250)
Storage condition:	Store in a dry place at room temperature

OTHER

AVAILABLE DOCUMENTS

Data Sheet	Certificate of Compliance
Certificate of Analysis	Safety Data Sheet (SDS)
Certificate of Sterility	Pyrogen Declaration
DNase/RNase	

Reach out to us for additional resources, if applicable to this product.

DECLARATION

CFIA	LABPLAS sampling bags are a solution that may be used in the CFIA Preventive Control Plan (PCP) for the seven principles of the HACCP system. The PCP is a Canadian federal initiative, under the Safe Food for Canadians Regulations (SFCR).
EU	The materials used to manufacture LABPLAS sampling bags meet, where applicable, the Eu No10/2011 standards for food contact with respect to particle migration.
DNase-free	This product is DNase-free. Sensitivity of 10-7 Kunitz units/µL
RNase-free	This product is RNase-free. Sensitivity of 10-9 Kunitz units/µL.
FDA	The plastic film used in the manufacture of the LABPLAS sampling bag meets the requirements of the Food and Drug Administration Regulation [21 CFR 177.1520 (b), (c)2.1, (c)3.1a, (c)3.2a, and 170.39, 174.5 (a), 178.2010 (b), 178.3297 (c), and 178.3860], provided that the sampling bag is not in contact with an alcoholic product and that the conditions of use comply with section C to G of table 2 of 21 CFR 176.170 (c).
Pyrogens	This product is non-pyrogenic at the endotoxin limit of 2.15 EU/device. Non-pyrogenicity is supported by endotoxin testing of randomly selected samples using the Limulus amebocyte lysate (LAL) gel assay according USP-NF <85> and <161> guidelines.
Sterile	Sterility is provided by dry heat during extrusion of the plastic at temperatures exceeding 428 F. The approach ensures a sterility assurance level (SAL) of 10-3. Continued process effectiveness is demonstrated through periodic sterility testing. Sterility testing follows the USP-NF <71> guideline.

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