

GENERAL INFORMATION

Category : Sterile sampling kits

Family : Sani-Stick

Lifespan: 5 years

TECHNICAL DESCRIPTION

This kit consists of a Twirl'em bag and a biocide-free sponge on a handle designed for vigorous sampling of various surfaces and reaching difficult areas.



Volume guide

¹Total volume: The total volume corresponds to the maximum capacity the bag can hold when it is filled to the brim, without the possibility of closing it.

²Closure volume: The closure volume corresponds to the maximum capacity the bag can hold when it is closed with a triple twist to ensure safe transport.

³Functional volume: The functional volume corresponds to the volume of diluent or sample according to the test requirements.

SPECIFIC INFORMATION

CONTAINER

Type of container :	Bag
Material :	Thermoplastic laminate
Color :	Transparent
Dimension :	152 x 267 mm / 6 x 10.5 in
Thickness :	3.5 mil.in / 0.0889 mm / 88.9 micron
Total volume ¹ :	1200 ml / 40 oz
Closure volume ² :	700 ml / 24 oz
Functional volume ³ :	100 ml / 3 oz
Printing type :	Writing area
Opening system :	Laser scoring line
Closing system :	Attachment with 2 round wires
Sterile :	Yes
End of product life :	Non-recyclable

INSERTION

Insertion type :	Polyurethane sponge
Description :	Hydrophilic polyurethane foam sponge. White colour. Dimensions (L x W): 3" x 1.25" (77 mm X 31.8 mm). Free from biocides.
Biodegradable :	Non-recyclable

INSERTION

Insertion type :	Rigid Handle
Description :	Rigid polypropylene handle assembled with sponge. Recyclable. Green color. Equipped with a contactless sponge release mechanism.
Biodegradable :	Recyclable

SOLUTION

Name :	NA
Type of solution :	NA
pH :	NA

OTHER

Sterile blue gloves are the perfect tool to complement your sampling bags and kits. The Blue polyethylene gloves are packed in individual pairs, and then irradiated to sterilize them. They are folded and packed optimally to maintain sterility.

AVAILABLE DOCUMENTS

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

PACKAGING INFORMATION

Outer box dimension : (W x D x H)	11.88 X 17.63 X 12.56
Box weight :	11.00 LB / 4.99 KG
Conditioning :	100 (10 x 10)
Storage condition :	Store in a dry place at room temperature

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).
FDA	The plastic film used in the manufacture of LABPLAS sampling bags complies with 21 CFR 177.1520 [(c) 3.2c] of the Food and Drug Administration. Its use is limited to temperatures below 212 F according to Table 2 of 21 CFR 176.170 (c).
FDA	The plastic film used in the manufacture of the LABPLAS sampling bag meets the requirements of 21 CFR 177.1520 of the Food and Drug Administration.
Biocide-free	Biocide-free material
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 amoebocyte lysate (LAL) gel assay according USP-NF <85> and <161> guidelines.
Sterile	Sterility is provided by irradiation. The sterilization dose ensuring a sterility assurance level (SAL) of 10-3 has been established according to ISO 11137.

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