# **KWS-22200**

# **TECHNICAL SHEET**



## **GENERAL INFORMATION**

Category: Sterile sampling kits

Family: Water sampling kit

Lifespan: 3 years

#### **TECHNICAL DESCRIPTION**

A kit consisting of a Twirl'em bag with safety tabs and printed area, and a sodium thiosulfate tablet that neutralizes biocides such as chlorine, iodine and oxidizing agents present in water.

# **SPECIFIC INFORMATION**

## **CONTAINER**

Type of container :	Bag
Material :	Polyethylene blend
Color :	Transparent
Dimension :	127 x 229 mm / 5 X 9 po
Thickness:	3 mil.in / 0.0762 mm / 76.2 micron
Total volume :	700 ml / 24 oz
Functional volume :	430 ml / 14 oz
Printing type :	Writing area
Opening system :	Perforated line
Closing system :	Attachment with 2 round wires
Sterile :	Yes
End of product life :	Recyclable

#### **INSERTION**

Insertion type :	NA
Description :	
Biodegradable :	NA



## **SOLUTION**

Name :	Sodium thiosulfate pill
Type of solution :	Sodium thiosulfate
pH:	N/A

## **OTHER**



# **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)	8.75 X 10.50 X 2.63
Box weight :	1.00 LB / 0.45 KG
Conditioning:	100 (1 x 100)
Storage condition :	Store in a dry place between 15°C and 25°C

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.
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.
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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