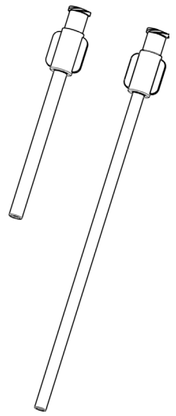
Nordson MEDICAL

Flexure™ Malleable Tip



Instructions for Use

www.nordsonmedical.com/eifu

Symbol Reference Key	
LATEX	Not made with natural rubber latex
REF	Catalog Number
	Manufacturer
	Caution
	Consult Instructions for Use
LOT	Lot Number
	Use By
STERILIZE	Do not resterilize
STERILE EO	Sterilized using Ethylene Oxide
	Do not use if the packaging is damaged
	Do Not Reuse
MD	Medical Device
	Single Sterile Barrier
MX	Country and Date of Manufacture
UDI	Unique Device Identification
MANUFACTURED IN MEXICO FOR:	
Manufactured in Mexico for:	
Malleable Tip	
Malleable Tip	

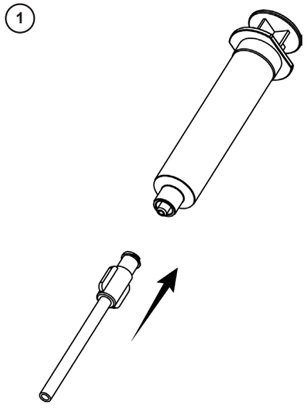
The symbol glossary is provided at
www.nordsonmedical.com/symbol glossary

MANUFACTURED IN MEXICO FOR:

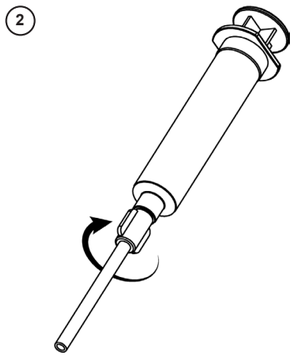
Micromedics, Inc.
1270 Eagan Industrial Road
St. Paul MN, 55121-1385 USA
651-452-1977
800-624-5662
info@nordsonmedical.com
www.nordsonmedical.com

M22065 Rev. B 2024-10-04

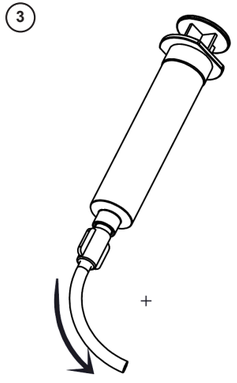
1



2



3



Flexure™ Malleable Tip

EN

Before using, read the following information:

DESCRIPTION / INTENDED USE:

The Malleable Tip is a sterile, single-use cannula intended for the application of materials in medical procedures.

CONTRAINDICATIONS:

None known.

WARNINGS / PRECAUTIONS:

- Do not use the device if the sterile packaging is damaged or unintentionally opened prior to use. Inspect the device before use. Do not use if the device appears to be damaged or defective and dispose per the Disposal instructions below.
- The device is intended for single use only. Do not resterilize. No effective cleaning process has been developed to prevent cross contamination. Contamination of a preprocessed device may lead to injury, illness or death of a patient.
- Do not cut or trim the applicator tip.
- Do not overtighten the Luer connector.
- There is a low probability that a delay in treatment, increased procedural time, mild tissue damage or local immunogenic response could occur if the device fails to function as intended.

INSTRUCTIONS FOR USE:

1. Remove applicator tip from sterile package using sterile technique and insert into luer lock connector. NOTE: Syringe not included.
2. Twist applicator tip firmly onto luer lock connector.
3. Applicator Tip is ready for use. NOTE: Tip may be angled to provide improved access.
4. Dispose of as Biohazard waste. Device will degrade if reprocessed. No effective cleaning process has been developed to prevent cross contamination. Contamination of a preprocessed device may lead to injury, illness or death of a patient.

DISPOSAL:

Dispose of as Biohazard waste. Device will degrade if reprocessed. No effective cleaning process has been developed to prevent cross contamination. Contamination of a preprocessed device may lead to injury, illness or death of a patient.

STORAGE CONDITIONS:

Sterilized products should be stored in a dry, clean environment, protected from direct sunlight, dust, pests, and extreme temperature and humidity.

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member state in which the user/patient is established.

United States only: 1-800-624-5662
Rx only For US Audiences Only

INTENTIONALLY LEFT BLANK

INTENTIONALLY LEFT BLANK

INTENTIONALLY LEFT BLANK