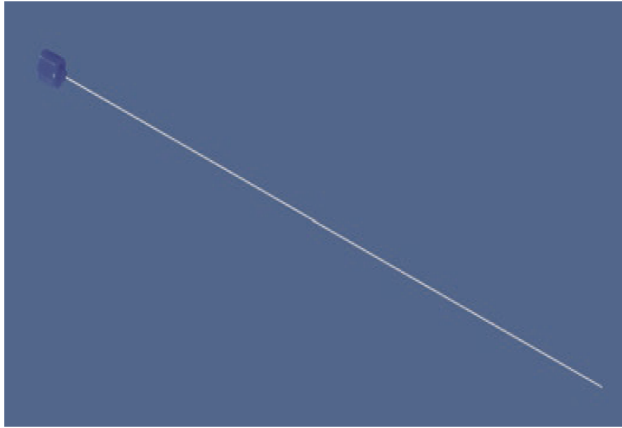




Not made with natural rubber latex

Description

The H/S Catheter Stylet is designed to use to gain ureteral access, to establish a tract, and to assist in the placement during gynecologic procedures.

Product Model

BA100

Precaution

1. Use caution not to force or over manipulate the H/S Catheter Stylet when gaining access.
2. When using H/S Catheter Stylet through a metal cannula, use caution as damage may occur to outer.
3. The H/S Catheter Stylet is not intended for PTCA use.

Caution

1. Sterile product unless package is opened or damaged.
2. Do not reuse for avoiding user may be infected by the microorganism.
3. This device is sold by or on the order of a physician.
4. If the sterile package is damaged or unintentionally opened before use, do not use.

Warnings

Dispose of medical waste properly in accordance with regulations after using.

Instructions for Use

1. Using aseptic technique, remove the H/S Catheter Stylet from its outer packaging and place in the sterile field.
2. Remove the H/S Catheter Stylet from the holder.
3. Adjust mandril position by advancing or withdrawing the mandril within the H/S Catheter Stylet to achieve the required degree of flexibility when using the H/S Catheter Stylet.
4. Advance the flexible end of the H/S Catheter Stylet into the HSG catheter and position to the desired location.

Note: The size and length of the device must be taken into consideration to ensure proper fit between H/S Catheter Stylet and device.

5. After H/S Catheter Stylet use complete, removed from patient and discard.

Glossary of Symbols

	Batch code
	Manufacturer
	Medical device
	Temperature limit 10-40°C
	Humidity limitation 30-80%RH
	Biological Risks
	Do not re-use
	Consult instructions for use
	Expiry Date (YYYY/MM/DD)
	Caution
	Do not use if package is damaged and consult instructions for use
	Single sterile barrier system and using ethylene oxide
	Catalogue number
	Not made with Natural Rubber Latex

Rx ONLY Federal (USA) law restricts this device to sale by or on the order of a physician or practitioner

Reporting of Serious Incidents

If a serious incident related to the device occurs, immediately report the incident to Bioteque Taiwan Ltd. and to the applicable competent authority having jurisdiction in their locale.



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