SWICKER DIGIT

Radiopaque Over-Finger Foam Sponge

Instructions for Use

DESCRIPTION:

SWICKER is a sterile foam sponge designed for single-patient use. **SWICKER** is a highly absorbent surgical sponge intended for use in a variety of surgical or medical procedures. **SWICKER** can be used in multiple surgical procedures and areas including patient prepping, open surgery, instrument padding, instrument cleaning, and wound care. **SWICKER DIGIT** is an over-the-finger configuration of the **SWICKER** product family. **SWICKER DIGIT** is designed to be used over a gloved finger.

<u>SWICKER</u> is packaged dry and can be used dry or hydrated. <u>SWICKER</u> is extremely soft, strong, and flexible. It resists tearing and is cellulose free. See below for <u>SWICKER</u> hydrating process.

SWICKER is sufficiently durable and tested to be used continuously during the same procedure and can be temporarily left in place during surgery. Suction can be applied to clear the **SWICKER** or wring it out. **SWICKER** contains radiopaque ink marker dots that are applied to the sponge.

INDICATIONS:

SWICKER is a sterile, single-use device indicated for use as a surgical sponge substitute. It can be used to protect soft tissues under surgical retractors and has significant strength and integrity to be used as a retraction device.

CONTRAINDICATIONS:

None known.

INSTRUCTIONS FOR USE:

- 1. Using aseptic techniques, peel open package and transfer SWICKER to sterile field/personnel.
- 2. **SWICKER** can be used dry to packing or for drying and cleaning a medical device.
- 3. To hydrate the **SWICKER**, submerge it and squeeze it a few times until it becomes completely saturated in a sterile bowl of sterile fluids.
- 4. Fold up or roll up to wring out excess solution into a trash receptacle prior to use.
- 5. SWICKER can be cleaned and reused continuously throughout the procedure by either wringing the blood or body fluids out in a bowl or by using suction to clear the sponge.
- 6. Wring out excess fluid prior to disposal. Dispose of contaminated devices in accordance with hospital policy and regulations for disposal of biohazard medical waste. Non-contaminated, unused devices may be disposed of per hospital guidelines.

PRECAUTIONS:

 SWICKER should be stored in its original sealed package at room temperature, away from direct sources of heat.

WARNINGS

- 1. Do not use **SWICKER** past the expiration date indicated on the label.
- 2. Do not use **SWICKER** if the sealed pouch is punctured, torn, or otherwise compromised.
- 3. Do not use **SWICKER** if the device is visibly torn, frayed, or damaged.
- 4. Do not re-sterilize. SWICKER is supplied sterile and is intended for single patient use only.
- 5. Do not leave **SWICKER** inside the body. It must be removed after use.

Table 1: Symbol Glossary

Symbol	Title (Reference)	Description
***	Manufacturer (5.1.1 ^[1])	Indicates the medical device manufacturer
>	Use-by date (5.1.4 ^[1])	Indicates the date after which the medical device is not to be used
LOT	Batch code (5.1.5 ^[1])	Indicates the manufacturer's batch code so that the batch or lot can be identified.
REF	Catalogue number (5.1.6 ^[1])	Indicates the manufacturer's catalogue number so that the medical device can be identified
STERILE EO	Sterilized using ethylene oxide (5.2.3 ^[1])	Indicates a medical device that has been sterilized using ethylene oxide
STERBAZZE	Do not resterilize (5.2.6 ^[1])	Indicates a medical device that is not to be resterilized
	Do not use if package is damaged and consult instructions for use (5.2.8 ^[1])	Indicates that a medical device that should not be used if the package has been damaged or opened and that the user should consult the instructions for use for additional information
(2)	Do not re-use (5.4.2 ^[1])	Indicates a medical device that is intended for one single use only.
elFU	Consult Instructions for Use (5.4.3 ^[1])	Indicates the need for the user to consult the instructions for use
	Single sterile barrier system (5.2.11 ^[1])	Indicates a single sterile barrier system
	Single sterile barrier system with protective packaging outside (5.2.14 ^[1])	Indicates a single sterile barrier system with protective packaging outside
UDI	Unique Device Identifier (5.7.10 ^[1])	Indicates a carrier that contains unique device identifier information
R only	Prescription only (21 CFR 801.109)	Caution: Federal law restricts this device to sale by or on the order of a physician

[1] ISO 15223-1:2021 Medical devices – Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.