Peri-Strips Dry®

With Veritas® Collagen Matrix • Circular Staple Line Reinforcement

INSTRUCTIONS FOR USE .................. 4
MODE D’EMPLOI ............................ 7
GEBRAUCHSANLEITUNG .................. 10
ISTRUZIONI PER L’USO .................. 14
INSTRUCCIONES DE USO ............... 17
GEBRUIKSINSTRUCTIES .................. 20
BETJENINGSVEJLEDNING .................. 23
BRUKSANVISNING ........................ 26
BRUKSANVISNINGER .................... 29
KULLANIM TALIMLARI .................... 32
ΟΔΗΓΙΕΣ ΧΡΗΣΗΣ .......................... 35
ΚΑΥΤΤΟΘΗΕΤ ......................... 39
INSTRUÇÕES DE UTILIZAÇÃO .......... 42
ISTRUCŢIUNI DE FOLOSIRE .......... 45
Peri-Strips Dry® with Veritas® Collagen Matrix Circular Staple Line Reinforcement

**Figures 1-6**

**FIGURE 1**
Circular Buttress Assembly

**FIGURE 2**
Cartridge Cone

**FIGURE 3**

**FIGURE 4**

**FIGURE 5**

**FIGURE 6**
Peri-Strips Dry® with Veritas® Collagen Matrix Circular Staple Line Reinforcement

*Figures 7-11*
DESCRIPTION:
Peri-Strips Dry® with Veritas® Collagen Matrix Circular Staple Line Reinforcement (PSD-V) is prepared from dehydrated bovine pericardium procured from cattle under 30 months of age in the United States.

The packaging contains a buttress assembly which includes two (2) circular buttresses, one for the anvil and one for the cartridge side of the stapler. Each buttress assembly has a coating of acrylic adhesive on one side for attachment to the stapler surfaces. The packaging also includes one cartridge cone to aid atraumatic advancement of the stapler into the surgical site. All components are packaged sterile.

INDICATIONS FOR USE:
PSD-V is intended for use as a prosthesis for the surgical repair of soft tissue deficiencies using surgical staplers when staple line reinforcement is needed. PSD-V can be used for reinforcement of staple lines during gastric, bariatric, and small bowel procedures.

CONTRAINDICATIONS:
The use of PSD-V is contraindicated in patients with known sensitivity to bovine or acrylic material.

ADVERSE REACTIONS:
As with any surgical procedure, adverse reactions are possible and include but are not limited to: infection, rejection, erosion, and allergic reaction.

WARNINGS:
Do not re-sterilize. Resterilization may cause changes to the tissue and negatively impact functionality of the device.

Do not use product if there is damage to the pouch or seals.

Ensure the staple line is completely covered with the buttress or inadequate coverage after firing may result.

SYMBOL DEFINITIONS:
- Store at a controlled room temperature.
- Keep away from heat. Do not use if heat indicator is red.
- Do not re-use
- Consult Instructions for Use
- Sterilized using Ethylene Oxide
- Do not use if the product sterilization barrier or its packaging is compromised.
- Product treated with sodium hydroxide.
- Product derived from USDA inspected cattle.
- Made in the U.S.A.
- CAUTION: Federal (USA) law restricts this device to sale by, or on the order of, a physician.
- Catalog number
- Use by date
- Lot number
- Compatible stapler models
- Authorized Representative in the European Community

CONTENT
- Rx Only
- Catalog number
- Use by date
- Lot number
- Compatible stapler models

Manufacturer

Authorized Representative in the European Community

SSI part number

SSI tracking number

SSI internal code

SSI internal code
PSD-V is not designed, sold, or intended for use except as indicated; doing so may result in surgical complications.
Synovis products differ; substitution of one product for another product may reduce product performance.
Do not use PSD-V if heat indicator has been activated.

CAUTIONS:
Do not get the anvil or cartridge buttress wet before applying, or the buttress may not adhere to the stapler properly.
Cartridge cones may be used for entry into the surgical site on the cartridge side of the stapler during endoscopic surgery. Before completion of surgery be sure to remove the cartridge cone from the surgical site.
Final tissue compression, including PSD-V, must meet the range specified by the stapler manufacturer. PSD-V increases the total thickness of the area stapled by 0.4 mm - 1.2 mm (0.016" - 0.048").
Follow Instructions for Use supplied by the stapler manufacturer. Do not use PSD-V contrary to the stapler manufacturer’s instructions.

PERI-STRIPS DRY WITH VERITAS COMPONENTS:
Each PSD-V package contains the following components (See Figure 1): a) one (1) pouch containing a circular buttress assembly which includes two (2) circular buttresses and b) one (1) pouch containing a cartridge cone.
Cartridge cone (See B): The cartridge cone is designed to protect the cartridge buttress during advancement of the stapler into the surgical site. There are two features to facilitate removal of the cone from the stapler and from the surgical site before completion of the surgery: 1. the indentations at the top of the cone are designed to allow the use of a forceps/hemostat to remove the cone from the stapler and/or surgical site, 2. the holes at the tip of the cartridge cone are designed to allow a suture to be placed in the cone to facilitate the removal of the cone from the surgical site.

INSTRUCTIONS FOR USE:
Each model of PSD-V has been designed specifically for the stapler models indicated on the label; verify that the correct model of PSD-V has been selected. Buttress Assemblies are identical and can be used on either the anvil or cartridge of the stapler.

Caution: Do not allow the buttresses, anvil, or cartridge surface to become wet. A wet buttress or a wet stapler surface may affect the ability of the buttress to adhere to the stapler surface.

Instructions for 21 mm or 25 mm circular staplers (Ethicon Endo-Surgery, Inc. and Covidien AutosutureTM):

A. Product Preparation
   A. Inspect the pouch(s). Do not use if any pouch is damaged or if the seal is not intact.
   B. Peel open the outer pouch of the buttress assembly and aseptically remove the inner pouch. The inner pouch may be placed in the sterile field.
   C. Peel open the pouch containing the cartridge cone and aseptically deliver the cone to the sterile field.

B1. Anvil Preparation
   D. Open inner pouch and remove the buttress assembly.
   E. Grasping the backing, peel away a single circular buttress (See Figure 2).
   F. Place the buttress over the anvil shaft with the adhesive side down towards the anvil surface (See Figure 3).
   G. Slide the buttress down the shaft of the anvil. The fit will be snug, use pressure to slide the buttress down the shaft.
   H. Press the buttress onto the surface of the anvil, ensuring that the buttress is centered and covers all staples and that there are no gaps between the buttress and the surface of the anvil (See Figure 4).
B2. Anvil Preparation for the Orvil Anvil Introducer

I. Open inner pouch and remove a buttress assembly.

J. Create a slit in one of the circular buttresses by cutting completely through one side of the buttress to the center hole (See Figure 5).

K. Grasp the backing and peel away the cut buttress from the backing (See Figure 6).

L. With the adhesive side down towards the anvil surface, slide the cut edge of the buttress over the shaft of the anvil (See Figure 7). Press the buttress onto the surface of the anvil, ensuring that the buttress is centered and covers all staples and that there are no gaps between the buttress and the surface of the anvil (See Figure 8). Ensure that there is no gap at the slit.

Note: Ensure proper alignment before applying pressure to the buttress.

C. Cartridge Preparation

A. Grasping the backing, peel away a single buttress. (See Figure 2).

B. Press the buttress onto the surface of the cartridge with the adhesive side down ensuring that the buttress is centered and covers all staples, and that there are no gaps between the buttress and the surface of the cartridge (See Figure 9).

D. Buttress Hydration

A. After applying buttresses to anvil and cartridge, immerse buttresses in sterile water or sterile saline for approximately 5 seconds.

C. Press the cone over the end of the stapler cartridge (See Figure 11).

IMPLANTING PSD-V INSTRUCTIONS FOR ALL STAPLER MODELS:

A. After inserting the stapler into the surgical site, remove the cartridge cone from the stapler by advancing the inner shaft or using a forceps/hemostat to grasp the tip of the cone. Note: If a suture has been placed in the cone, do not grasp or apply tension to the suture while removing the cone.

B. Follow the stapler manufacturer Instructions For Use to fire and remove the stapler. After firing, removal of the circular stapler through the buttressed anastomosis may be more difficult due to the reinforcement of the staple line.

C. Before completion of the surgery, remove cartridge cone from the surgical site either by pulling on the suture placed through the tip of the cartridge cone or by grasping with a forceps/hemostat.

DISCLAIMER OF WARRANTIES

Synovis Surgical Innovations (SSI), a division of Synovis Life Technologies, Inc., warrants that reasonable care has been used in the manufacture of this device. This warranty is exclusive and in lieu of all other warranties whether expressed, implied, written, or oral, including, but not limited to, any implied warranties of merchantability or fitness. As a result of biological differences in individuals, no product is 100% effective under all circumstances. Because of this fact and since SSI has no control over the conditions under which the device is used, diagnosis of the patient, methods of administration or its handling after it leaves its possession, SSI does not warrant either a good effect or against an ill effect following its use. The manufacturer shall not be liable for any incidental or consequential loss, damage or expense arising directly or indirectly from the use of this device. SSI will replace any device, which is defective at the time of shipment. No representative of SSI may change any of the foregoing or assume any additional liability or responsibility in connection with this device.
Authorized Representative in the European Community:
AR-MED
Runnymede Malthouse Business Centre
Egham, Surrey TW20 9BD
United Kingdom

Peri-Strips Dry and Veritas are registered trademarks of Synovis Life Technologies, Inc., (US Patent #6,312,474; 6,652,594; Additional Patents Pending)