ORDER HEMOBLAST™ BELLOWS

HIGH-PERFORMANCE HEMOSTASIS MADE SIMPLE.

HEMOBLAST™ Bellows is supplied as a preloaded bellows that:

- Contains 1.65 g of powder, including a maximum of 1,500 units of thrombin
- Includes a 10 cm nozzle extension to assist with application where a slightly longer tip is desired to reach the target bleeding site in open procedures
- Is biocompatible and non-pyrogenic

MORE VERSATILITY FOR MORE APPLICATIONS

The HEMOBLAST™ Bellows Laparoscopic Applicator is:

- Intended for delivering HEMOBLAST™ Bellows hemostatic powder to bleeding surgical sites
- Compatible with a 5 mm diameter or larger trocar
- Sold separately from HEMOBLAST™ Bellows*

*Please see the HEMOBLAST™ Bellows and HEMOBLAST™ Bellows Laparoscopic Applicator Instructions for Use for complete information on the hemostatic agent.

PART NUMBER	DESCRIPTION	MINIMUM ORDER QTY
BQF02-US	1 ea. HEMOBLAST™ Bellows and 1 ea. 10 cm Nozzle Extension	1 case/12 ea.
LAP01-US	1 ea. Individually Packaged Laparoscopic Applicator	1 case/12 ea.

REQUEST A CLINICAL EVALUATION OR DEMONSTRATION:

1-877-GO-DILON (1-877-403-4566)

Hemoblast.com/us/evaluation

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orders@dilon.com

Caution: Federal law restricts this device to sale on or by the order of a physician.

Important Risk Information for HEMOBLAST™ Bellows: Do not inject HEMOBLAST™ Bellows into a vessel or tissue. There is a risk of allergic-anaphylactoid reaction and/or thromboembolic events, which may be life-threatening. Do not apply HEMOBLAST™ Bellows in the absence of active blood flow, e.g., while the vessel is clamped or bypassed. Extensive intravascular clotting and even death may result. Do not administer to patients with known allergies or hypersensitivity to materials of porcine or bovine origin. Because HEMOBLAST™ Bellows is made from human blood, it may carry a risk of transmitting infectious agents, e.g., viruses, and theoretically, the Creutzfeldt-Jakob disease (CJD) agent. HEMOBLAST™ Bellows contains chondroitin sulfate from bovine origin which is associated with a remote risk for Transmissible Spongiform Encephalopathies (TSE), which has been minimized in accordance with regulatory guidelines by a manufacturing process with demonstrated TSE inactivation capacity. When applied to a bleeding site, HEMOBLAST™ Bellows swells up to 60% within about 5 minutes. Do not attempt to trim the applicator tip. HEMOBLAST™ Bellows should not be used at the site of a valve replacement or repair as valvular dysfunction could occur. HEMOBLAST™ Bellows should not be applied at the site of a synthetic graft or patch implant due to potential for decreased effectiveness. The product should not be in contact with circulating cerebrospinal fluid (CSF). The material has not been tested on children and pregnant or lactating women. Rx Only. For safe and proper use of this device, refer to the full Instructions for Use.

Important Risk Information for HEMOBLAST™ Bellows Laparoscopic Applicator: Do not use the applicator to manipulate or retract organs or tissue. The product must be manipulated and used by qualified personnel according to the general principles of sterility and pre-medication. The tip of the laparoscopic applicator must be directly visualized at all times to minimize potential for unintended contact with tissue, organs, or blood as well as any possible unintended spillage of residual powder. Rx Only: For safe and proper use of this device, please refer to full device Instructions for Use.

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