PuraBond®
Absorbable Haemostat

No waiting time after declamping in cardiac surgery*

Pre-filled syringe, **ready to use**, NO preparation required

No clotting in the syringe

Achieves haemostasis **in seconds**(2)

* when there is oozing from needles holes in anastomoses and haemostasis by ligation or standard means is insufficient or impractical.

1. PuraBond IFU-004 Rev 1.1.
2. Morshuis M, Final Clinical Study Report: A Single-Centre, Single Arm Post-Market Clinical Follow-Up to Confirm the Safety and Performance of PuraStat Absorbable Haemostatic Material for the Management of Bleeding After Left Ventricular Assist Device (LVAD) Implantation, BR CV 003 EU EN v1 2019 09 13
3. Adapted from: https://stock.adobe.com/uk/images/female-nurse-and-male-doctor/344055235

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Application Guidance

Oozing bleed or resected area[1]

1

Step 1 - Clear
Remove as much blood and fluid as possible.

2

Step 2 - Direct
Apply PuraBond at the base of the lesion, as close as possible to the source of bleeding.

3

Step 3 - Apply
Apply PuraBond in an even layer until the whole lesion is covered.

4

Step 4 - Wait
Do not disturb PuraBond until sufficient time is allowed for haemostasis.

If Necessary, Re-Application Guidance

5

Step 5 - Direct
Place applicator through the previous PuraBond layer directly onto source of bleeding.

6

Step 6 - Re-apply
Apply PuraBond at the base of the lesion, as close as possible to the source of bleeding.

7

Step 7 - Wait
Do not disturb PuraBond until sufficient time is allowed for haemostasis.

After haemostasis is confirmed, excess PuraBond may be left in place

PuraBond is available in quantities of
1 mL (621-201)
3 mL (621-202)
5 mL (621-203)

PuraBond is indicated for haemostasis in the following situations encountered during surgery, when haemostasis by ligation or standard means is insufficient or impractical[1]:

- Oozing from vascular anastomoses
- Bleeding from small blood vessels and oozing from capillaries of the parenchyma of solid organs

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(1) PuraBond IFU-004 Rev 1.1.

PuraBond is a class III medical device, CE marked according to European council directive 93/42/EEC on medical devices and its relatives

CE 2797 BSI