Hemopatch
Sealing Haemostat

Application Steps – as easy as 1, 2, 3...

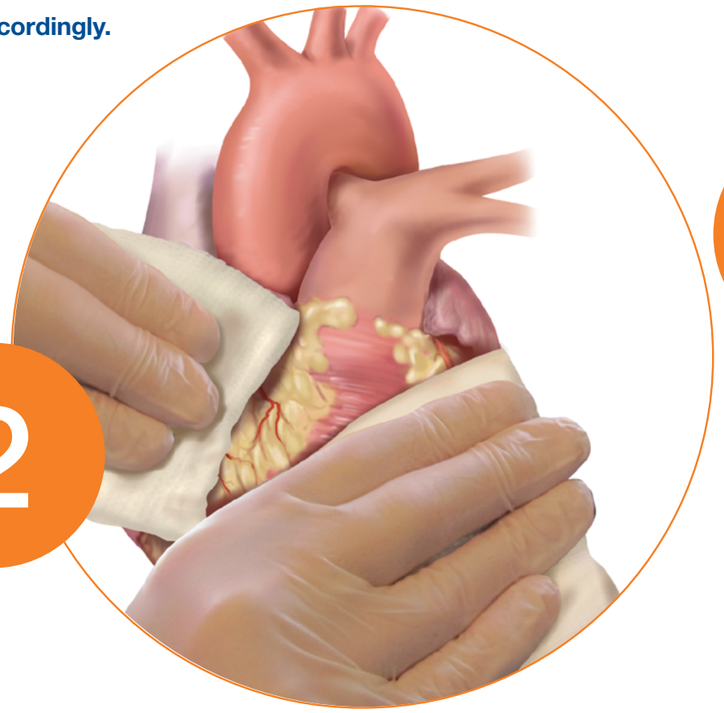
HEMOPATCH is intended as a haemostatic device and surgical sealant for procedures in which control of bleeding or leakage of other body fluids or air by conventional surgical techniques is either ineffective or impractical. HEMOPATCH may be used to close dural defects following traumatic injury, excision, retraction or shrinkage of the dura mater.[1]

HEMOPATCH comes ready to use in sterile packages and must be handled accordingly.
Use only undamaged packages, single use only, do not re-sterilise.

1

Apply HEMOPATCH **dry** with blue dots facing **up** and the white non-marked side in contact with the bleeding surface

2

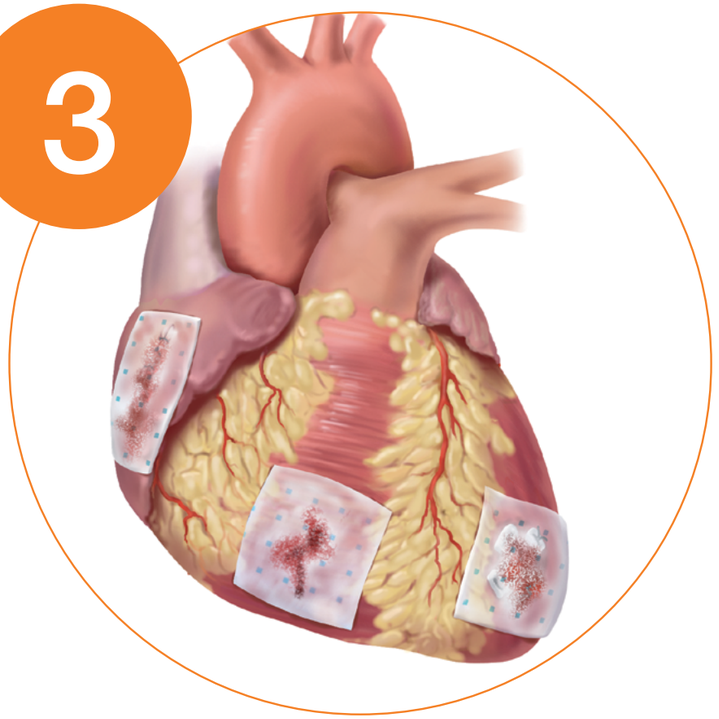


Use a **dry gauze** and approximate HEMOPATCH with gentle uniform pressure for 2 minutes

Improved adherence is observed when HEMOPATCH is in direct contact with wound fluid such as blood or lymphatic fluid

In the absence of such wound fluids, sodium bicarbonate solution (concentration between 4.2% to 8.4%) may be used in conjunction with HEMOPATCH application

3



Confirm haemostasis has been achieved and leave HEMOPATCH in situ; **avoid disrupting the clot-formation**

For full product information please refer to the instructions for use

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Ordering information	Size	Units per box	Order Number
HEMOPATCH Small	2.7 x 2.7 cm	5	1506257
HEMOPATCH Medium	4.5 x 4.5 cm	3	1506256
HEMOPATCH Large	4.5 x 9.0 cm	3	1506253

Any medical device product quality complaints (including medical device adverse incidents) relating to Baxter Country Quality Assurance Team:
In the UK on +44 (0)1604704603, or by email to UK_SHS_QA_Complaints@Baxter.com. In Ireland on +353 (0)12065500 or by email to SHS_Complaints_Dublin@Baxter.com.
Alternatively please report directly to your Baxter Representative, who will take the details and forward to the Baxter Country Quality Assurance Team.
Medical device adverse incidents should also be reported: In the UK to the MHRA.
Reporting forms and information can be found at: www.mhra.gov.uk/safetyinformation/reportingsafetyproblems/index.htm
In Ireland to the HPRA. Reporting forms and information can be found at: http://www.hpra.ie/homepage/about-us/report-an-issue

1. HEMOPATCH Sealing Hemostat Instructions for Use. Zurich, Switzerland: HEMOPATCH IFU M000483. March 2017.

For detailed information please contact your local representative

Baxter

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