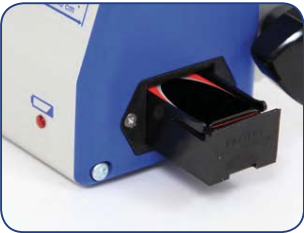




# EASYSpray Quick Reference Guide (for open wound surgery)





## Instructions for Circulating Nurse | EASYSpray Pressure Regulator


- 

Insert 9V battery into the EASYSpray pressure regulator device.
- 

Connect EASYSpray device to IV pole or trolley rail using the clamps on the back of the device.
- 


Use suitable connection tube to connect the EASYSpray device to medical air (ranging 3.5 – 7 bar / 51 – 100 psi).
- 


Connect Spray Set filters to EASYSpray device. Connect the blue filter (pressure line) to the blue female luer connector and the clear filter (sensor line) to the male luer connector.
- 


Turn the on/off switch on the front side of the EASYSpray to the ON (I) position.
- 


Check the gauge on the EASYSpray device for the appropriate pressure range of 1.5-2.0 bars (21.5-28.5 psi). Adjust pressure setting by turning the grey pressure control knob.


## Instructions for Scrub Nurse | Spray Set


- 

Prepare TISSEEL Fibrin Sealant according to the instructions in the package insert.
- 

Firmly attach the spray head to the nozzle of the syringes.
- 


Fasten the pull strap to the double syringe system to assure the spray head is tightly secured.
- 


Fit the connection tube of the spray set to the luer-lock connector on the underside of the spray head.
- 


Attach the clip (on the end of the sensor line) by sliding it into the grooves located on the top of the syringe plunger.
- 

Pass the assembled applicator to the surgeon for spray application. Pass the end of the connection tube with the sterile filters to the circulating nurse.

## Instructions for Surgeon

- 

Confirm (verbally) the actual pressure with theatre personnel.
- 

Spray from a distance of 10 – 15 cm for optimum results.
- 

To activate the flow of gas occlude the opening in the clip center with thumb. To begin application, gently depress the syringe plunger.

The use of TISSEEL Fibrin Sealant is restricted to experienced surgeons who have been trained in the use of TISSEEL. In order to ensure optimal safe use of TISSEEL by spray application, apply a minimum spray distance of 10 cm and a maximum spray pressure of 2 bar to minimise the potential risk of air or gas embolism, tissue rupture, or air or gas entrapment with compression.

The EASYSpray device will continue to emit gas for a brief period after the thumb is removed from the clip/plunger. This delay helps to avoid clogging of the spray head.

Please see the TISSEEL SmPC provided together with this material.

**Caution:** Any application of pressurised gas may be associated with a potential risk of air or gas embolism, tissue rupture or air or gas entrapment with compression, which may be life threatening. Be sure to take appropriate measures to address these risks by observing the recommended minimum spraying distance and the maximum pressure provided in the appropriate spray set instructions for use.

# EASYSpray Quick Reference Guide



## CAUTION BOX

Any application of pressurized gas may be associated with a potential risk of air or gas embolism, tissue rupture or air or gas entrapment with compression, which may be life threatening if the product is sprayed incorrectly

### Precautions

For TISSEEL/TISSUCOL Fibrin Sealant

- In open-wound surgery: when applying sprayable fibrin solutions for sealant using a pressure regulator device, the maximum pressure should be 2.0 bar (28.5 psi). The product should be sprayed at a distance at least 10 cm from the tissue surface.
- Spray application should only be used if it is possible to accurately judge the spray distance as recommended by the manufacturer. Do not spray closer than the recommended distances.
- Prior to applying sprayable fibrin solutions for sealant, the surface area of the wound should only be dried using standard techniques (eg, intermittent application of compresses, swabs, use of suction devices).
- Blood pressure, pulse rate, oxygen saturation and end tidal CO<sub>2</sub> should be monitored closely when spraying fibrin solutions for sealant using a pressure regulator device, because of the possibility of occurrence of air or gas embolism.
- Regulators should be used in line with manufacturer recommendations and the SmPC and Instruction for Use.

In order to ensure optimal safe use of TISSEEL by spray application the following recommendations should be followed:

Recommended pressure, distance and devices for spray application of TISSEEL					
Surgery	Spray set to be used	Applicator tips to be used	Pressure regulator to be used	Recommended distance from target tissue	Recommended spray pressure
Open wound	TISSEEL / ARTISS Spray Set	n.a.	EASYSpray	10-15 cm	1.5-2.0 bar (21.5-28.5 psi).
	TISSEEL / ARTISS Spray Set 10 pack	n.a.	EASYSpray		
	TISSEEL/TISSUCOL Spray Set	n.a.	EASYSpray		

When spraying the TISSEEL, changes in blood pressure, pulse, oxygen saturation and end tidal CO<sub>2</sub> should be monitored because of the possibility of occurrence of air or gas embolism.

### WARNINGS AND PRECAUTIONS

Caution: Pressurized gas supply must be between 3.5 and 7 bar (51-100 psi).

Caution: DO NOT connect to an oxygen source. Failure to follow these instructions can lead to explosion, which could result in serious injury or death.

Prior to operation make sure that the pressure range of the device is adjusted to the range specified in the appropriate spray set instructions for use.

Caution: Spray application requires appropriate safety measures as described in the pertinent Summary of Product Characteristics and Instructions for Use of the spray devices to make sure that the risks mentioned in this reference guide will be avoided

NOTE: Do not use with rechargeable batteries. The use of accessories of other manufacturers is not permitted.

<http://www.baxterspraysafety.com/index.html>

For detailed information please contact your local representative.

**Baxter**

Baxter Healthcare Ltd. Wallingford  
Road, Compton Newbury,  
Berkshire, RG20 7QW

In some countries TISSEEL Fibrin Sealant is licensed under the trademark of TISSUCOL Fibrin Sealant.

Baxter, Tisseel, Tissucol, and Easyspray are trademarks of Baxter International Inc., its subsidiaries or affiliates.

UK/128/19-0002a July 2019

### PRESCRIBING INFORMATION - TISSEEL Ready to use Solutions for Sealant

(Please consult the Summary of Product Characteristics before prescribing)

**Composition:** prefilled double chamber syringe containing deep frozen Sealer Protein Solution (with aprotinin) and Thrombin Solution (with Calcium Chloride Dihydrate). Sealer Protein Solution contains 91mg/ml human fibrinogen (clottable protein) and 3000 KIU/ml aprotinin. Thrombin Solution contains 500 IU/ml human thrombin and 40µmol per ml calcium chloride. Presentations of 1, 2 or 5ml in each chamber resulting in total volume of 2ml, 4ml or 10ml of sealant. **Indications:** Supportive treatment where standard surgical techniques are insufficient, for improvement of haemostasis, as a tissue glue to promote adhesion, sealing or as suture support, in gastrointestinal anastomoses, in neurosurgery where contact with cerebrospinal fluid or dura mater may occur and for mesh fixation in hernia repair, as an alternative or adjunct to sutures or staples. **Dosage and Route:** For epilepsial (topical) use only. Use of TISSEEL is restricted to experienced surgeons who have been trained in the use of TISSEEL. A thin layer is applied under direct vision to the tissue surface where required. Dose depends on the indication, application method and number of applications. For tissue adherence, it is recommended that the initial application cover the entire intended application area. As a guideline for the gluing of surfaces, 1 pack of TISSEEL 2 ml will be sufficient for an area of at least 10 cm<sup>2</sup>. Tissue surface should be as dry as possible before application. Do not use pressurized air or gas for drying the site. Application can be repeated if necessary but avoid reapplication of TISSEEL to pre-existing polymerized TISSEEL. Apply by drops or spray as needed depending on indication. **Side effects:** See Summary of Product Characteristics for detail. Postoperative wound infections. Fibrin degradation products increased. Hypersensitivity/anaphylactic reactions, anaphylactic shock, paresthesia, bronchospasm, wheezing, pruritus, erythema. Sensory disturbance. Bradycardia, tachycardia. Axillary vein thrombosis, hypotension, haematoma, embolism arterial, air embolism, cerebral artery embolism, cerebral infarction. Dyspnoea. Nausea, Intestinal obstruction. Rash, urticaria, impaired healing. Pain in an extremity. Procedural pain, pain, increased body temperature, flushing, oedema. Seroma, angioedema. **Class reaction:** Air or gas embolism, see Precautions. **Precautions:** Life threatening thromboembolic complications may occur if unintentionally applied intravascularly. Apply with care in coronary artery bypass surgery due to increased risk of inadvertent intravascular application. Must not be injected into highly vascularized tissue, such as nasal mucosa. When spraying TISSEEL, changes in blood pressure, pulse, oxygen saturation and end tidal CO<sub>2</sub> should be monitored because of the possibility of occurrence of air or gas embolism. Air or gas embolism, tissue rupture, or gas entrapment with compression, which may be life-threatening, have occurred with the use of spray devices employing a pressure regulator to administer fibrin sealant at higher than recommended pressures and in close proximity to the tissue surface. When applying by spray, follow the instructions provided with the spray device, with particular reference to gas pressure and distance from the tissue surface. Do not administer with spray devices in enclosed body areas. Take risk of compressive complications into account when applying in confined spaces. Use with caution in patients with prior exposure to aprotinin. Caution in patients with bovine protein allergies. Infectious diseases due to the transmission of infective agents cannot be totally excluded. Use of Tisseel and batch number should be recorded in patient's notes. Excessive clot thickness may negatively interfere with product efficacy and the healing process. Oxidised cellulose-containing preparations should not be used with TISSEEL. **Contraindications:** Do not apply intravascularly. Hypersensitivity to active substances or other components. Not for the treatment of active or spurting arterial or venous bleeding. Not for replacement of skin sutures intended to close surgical wounds. Interactions: Avoid solutions containing alcohol, iodine and heavy metals. **Overdose:** Not reported. **Legal category:** POM. **Basic NHS price:** 2ml - £97.50; 4ml - £195.00; 10ml - £443.75. **Marketing Authorisation Number and Holder:** PL 00116/0627 Baxter Healthcare Limited, Caxton Way, Thetford, Norfolk IP24 3SE, UK. **Date of preparation:** Jan 2019

Adverse Events and any drug or medical device product quality complaints (including suspected defective medicines or medical device adverse incidents) should be reported. Reporting forms and information can be found at [www.mhra.gov.uk/yellowcard](http://www.mhra.gov.uk/yellowcard). Adverse Events should also be reported to Baxter Healthcare Ltd, by email ([vigilanceuk@baxter.com](mailto:vigilanceuk@baxter.com)) or by phone (+44(0)1635206360). Drug or medical device product quality complaints relating to Baxter products can be reported directly to Baxter Healthcare Ltd by email ([UK\\_SHS\\_QA\\_Complaints@baxter.com](mailto:UK_SHS_QA_Complaints@baxter.com)) or by phone (+44 (0) 1604 704603).