Made in Italy by

EMODIAL

CE
0373

VALIANT MEDICAL

EXIT-PRO® AG

Silver Dressing

VALIANT MEDICAL

www.valiantmedical.co.uk
+44(0)330 303 9090
info@valiantmedical.co.uk

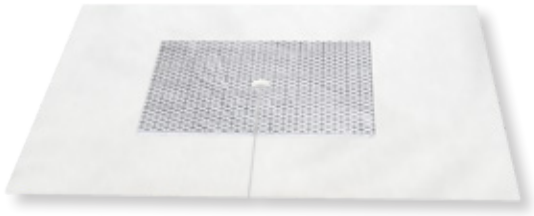
VM-EXITPRO AG-BROCHURE-04/21

EXIT-PRO® AG

DESCRIPTION
Non adherent silver ions based dressing set on breathable hypoallergenic adhesives:
- nonwoven tissue plaster;
- waterproof polyurethane membrane.

INDICATIONS
Specific for the exit site dressing.
To be used for compression and absorption of the exudates.

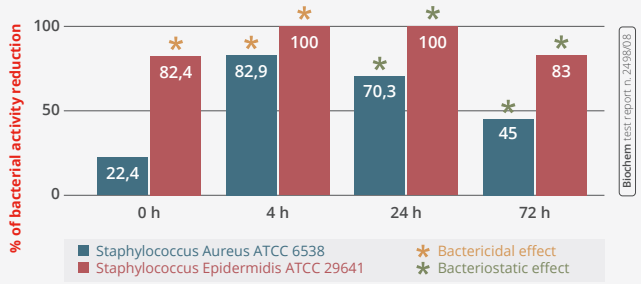
PURPOSE
- Immediate and long term antibacterial action. It helps to reduce the risk of local infection of the exit-site, of the tunnel or systemic (catheter-correlated sepsis) related to the use of intravascular devices.
- It helps to prevent the irritation caused by the mechanical rubbing of the catheter over the skin.
- The waterproof adhesive model protects the exit-site from contact with water during patient's hygiene.
- Patients with hyperhidrosis or hyperthermia are recommended to use the model with non woven adhesive.



REFERENCES
Data and technical specifications can be changed without notice.

CODE	PLASTER (cm)		PAD (cm)	MODEL (cm)	PCS / PACK	PACKS / BOX
	NW	PU				
S0810Y/AG	10 x 7		5 x 4	CUT AND HOLE ø 0.5	1	100
S0810YH/AG		10 x 8	5 x 4		1	100
S0810F/AG	10 x 7		5 x 4	CUT AND HOLE ø 0.5	1	100
S0810FH/AG		10 x 8	5 x 4		1	100
S0805F/AG	10 x 7		5 x 4	CUT AND HOLE ø 0.4	1	100
S0805FH/AG		10 x 8	5 x 4		1	100
S0808/AG	8 x 3		1.8 x 3.5	Y CUT	2	100

Antibacterial Action



METHOD OF ANALYSIS
A standardised bacterial culture (minimum starting concentration of 1x106 cfu/ml) is placed onto the sample (pad with silver) and control (pad without silver). The count of the bacterial colonies grown is made immediately after sowing, after 4, 24 and 72 hours of incubation period. The results are expressed as percentage bacterial reduction in **EXIT-PRO® AG** sample vs. control.

DIRECTIONS FOR USE
1. Disinfect the Exit-Site (avoiding iodine based disinfectants) and let it completely dry.
2. Put **EXIT-PRO® AG** at the catheter Exit-Site and hold in place by pressing on the adhesive.
3. Change **EXIT-PRO® AG** as per hospital protocol, preferably once per procedure, or whenever it gets wet, dirty or a site inspection is needed.
4. Model B with a waterproof polyurethane membrane protects the fistula from contact with water during patient's hygiene.

⚠ WARNINGS
- Do not use in patients with hypersensitivity to silver.
- Do not use during examinations such as X-rays, ultrasound, diathermy, magnetic resonance or radiation treatments.
- It does not replace the appropriate systemic therapies required in case of clinical infections.
- As usual with adhesive devices, remove it with extreme caution.

CLASSIFICATION	CLASS III MEDICAL DEVICE (EUROPEAN COMMUNITY)
BIOCOMPATIBLE	ACCORDING TO UNI EN ISO 10993 STANDARDS
STERILIZATION	GAMMA RAYS
EXPIRATION	3 YEARS
⚠ ②	SINGLE-USE