

Made in Italy by



EXIT-PAD[®]

S I L V E R **AG**

Silver dressing

SILVER
SILVER



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EXIT-PAD®

AG

S I L V E R

PAT. PEND.

Description:

Non adherent silver dressing.

Indications:

Specific for the exit site dressing.

To be used for the compression and the absorption of exudates.

Purpose:

1. The silver layer has an immediate and long term antibacterial action helping 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.
2. It helps to prevent the irritation caused by the mechanical rubbing of the catheter over the skin.



REFERENCES

Data and technical specification can be changed without notice

CODE	NON ADHERENT SILVER DRESSING		
	PAD measure mm	Model	Qty blister pack
S0903/AG	35X45	With cut and hole ø 4 mm	1
S0904/AG	35X45	With cut and hole ø 5 mm	1
S0925/AG	35X22	With cut and hole ø 4 mm	2
S0905/AG	50X40	With Y cut	1

Sperimental evidences

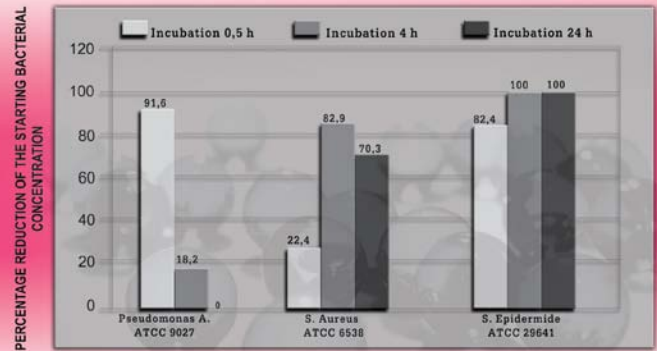
The antibacterial action **EXIT-PAD-Ag pads** has been tested.

The results are shown in the down below diagram.

Biochem test report n°2498/08

EXIT-PAD®

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Method of the analysis

A standardised bacterial culture (minimum starting concentration of 1×10^6 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 (0,5 h), after 4 h and 24 h incubation period. The results are expressed as percent bacterial reduction in **EXIT PAD-Ag** sample vs.control.



Direction for use

- 1 Disinfect the Exit-Site (avoiding iodine based disinfectants) and let it completely dry.
- 2 Put Exit-Pag-Ag® at the catheter Exit-Site and hold in place with plaster or a sterile semipermeable breathable dressing.
- 3 Change Exit-Pag-Ag® as per hospital protocol, preferably once per procedure, or whenever it gets wet, dirty or a site inspection is needed.

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

Classification: Class III medical device
Biocompatibility: The material in contact with the skin has passed the biocompatibility tests according to UNI EN ISO 10993 standards

Sterilisation: γ - ray
Expiry: 3 years