Made in Italy by

EMODIAL

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0373

VALIANT MEDICAL

NOVA SOVAN® AG

Silver Fistula Plaster

VALIANT MEDICAL

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VM-SOVAN AG-BROCHURE-04/21

NOVA SOVAN® AG

DESCRIPTION
Non adherent silver pad set on breathable hypoallergenic adhesives:
- non woven tissue;
- waterproof polyurethane membrane.

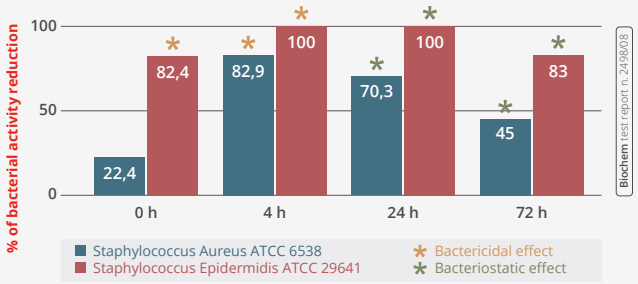
INDICATIONS
Compressive haemostatic action, specific for post-dialysis arterovenous access care to be used after the removal of fistula needles.
Also suitable for patients who require intravenous treatment and blood transfusions. Particularly suitable for the dressing of arteriovenous fistulas managed with the method of button-hole.

PURPOSE
- Reduces the risk of the arteriovenous fistula infection thanks to the presence of the silver based film with long-term and immediate antibacterial action.
- Reduces signs of inflammation associated with repeated venipuncture of the AVF.
- Prevents eschar formation and makes less traumatic for the patient the next venipuncture.
- The waterproof adhesive model protects the fistula from contact with water during patient's hygiene.

REFERENCES
Data and technical specifications can be changed without notice.

CODE	PLASTER (cm)		PAD (cm)	PCS / PACK	PACKS / BOX
	NW	PU			
S0079-1/AG	10 x 5		2.5 x 2.5	1	40
S0079/AG	10 x 5		2.5 x 2.5	2	30
S0079-1H/AG		10 x 5	2.5 x 2.5	1	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 NOVA SOVAN® AG sample vs. control.

DIRECTIONS FOR USE
1. Remove the first half of the protective film from the plaster or waterproof adhesive.
2. Put the pad over the needle site.
3. Remove the second half of the protective film and fix the plaster or the waterproof adhesive to the skin.
4. Model B with a waterproof polyurethane membrane protects the fistula from contact with water during patient's hygiene.

⚠ WARNINGS
- Avoid iodine based disinfectants.
- 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
⚠ ⊘	SINGLE-USE