Aniosyme Foam

Instrument Cleaning

Pre-treatment of surgical instrumentation in foam

Foam spray for cleaning of medical and surgical instruments. Ensures optimised dissolving of contamination on instruments with longer standing time (up to 72 hours) or before machine processing.

Aniosyme Foam is:

- **EASY** - Simple to use
- **FAST** - Prevents drying out of contamination
- **EFFECTIVE** - Stable foam with good wetting

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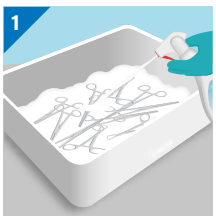
Spray

Instructions for use



Ready-to-use solution. Beforehand, spray the bottom of the soaking tray with Aniosyme Foam and place instruments collected.

Re-cover all the instruments with foam.

Cover the tray with the lid.

Thoroughly rinse treated medical devices with tap water to remove all traces of product and organic soiling.

Not suitable for instruments made of zinc or iron.

Packaging and ordering information

Product	Packaging unit	Order code
Aniosyme Foam	12 x 750 mL bottles	2380544

Application area

Pre-treatment of surgical instrumentation in foam: during a dry mode of transport or during storage of soiled instruments before machine or manual washing. Maintains an adequate level of humidity to avoid blood and other organic materials drying.

Instructions for use

Ready-to-use solution.

1. Beforehand, spray the bottom of the soaking tray with Aniosyme Foam and place instruments collected.
2. Re-cover all the instruments with foam.
3. Cover the tray with the lid.
4. Thoroughly rinse treated medical devices with tap water to remove all traces of product and organic soiling.

Not suitable for instruments made of zinc or iron.

Composition

Non-ionic surfactants, enzymatic complex (protease, lipase, and amylase), chelating agent, excipients.

Storage

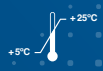
Dangerous – respect the precautions for use (Drawn up according to the European rules in force regarding the classification and labelling of chemical products).
Storage: between +5°C and +25°C.

Certificate

Scientific dossier (efficacy, compatibility, etc.) available on request.

Use medical devices safely

Always read the label and product information before use. Any serious incident related to the medical device must be reported to the manufacturer and the competent authority of the Member State.



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LABORATOIRES ANIOS
1, Rue de l'Espoir
59260 Lezennes
France

Distributed by:

ECOLAB DEUTSCHLAND GmbH
Ecolab-Allee 1
40789 Monheim am Rhein
Germany

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