

Summary of Product Characteristics

1 NAME OF THE VETERINARY MEDICINAL PRODUCT

Aluspray 4% w/w Cutaneous Spray, Powder

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Active Substance

Aluminium powder 4.0% w/w

Excipients

For a full list of excipients see section 6.1.

3 PHARMACEUTICAL FORM

Cutaneous spray, powder.

A grey, oily powder.

4 CLINICAL PARTICULARS

4.1 Target Species

Cattle, horses, swine, poultry, sheep, dogs and cats.

4.2 Indications for use, specifying the target species

For the protection and healing of wounds.

4.3 Contraindications

Do not use in case of hypersensitivity to the active substance or to any of the excipients.

Do not use in animals suffering from severe renal impairment.

Do not use to treat teat injuries in lactating animals producing milk for human consumption.

4.4 Special warnings for each target species

None.

4.5 Special precautions for use

Special precautions for use in animals

The product should be used unaltered from the original container.

In case of blockage of the spray nozzle, immerse it in hot water.

Special precautions to be taken by the person administering the veterinary medicinal product to animals

This product stains: avoid contact with hands and clothes.

Wear gloves when administering the product, wash hands after use.

Treat animals outdoor or in a well ventilated area.

Avoid inhalation. Do not swallow.

Pressurised container.

Flammable.

Do not spray on or near a naked flame or any incandescent material.

4.6 Adverse reactions (frequency and seriousness)

None.

4.7 Use during pregnancy, lactation or lay

Pregnancy:

Can be used during pregnancy.

Lactation:

Do not use to treat teat injuries in lactating animals producing milk for human consumption.

4.8 Interaction with other medicinal products and other forms of interactions

None.

4.9 Amounts to be administered and administration route

Shake well before use.

A superficial application onto wounds once or twice daily, to produce a fine coating of the powder. For optimal use, wounds should be cleaned, disinfected and sutured if necessary before the application of Aluspray.

4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary

None.

4.11 Withdrawal period(s)

Zero days.

Not authorised for use in lactating animals producing milk for human consumption.

5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES

Pharmacotherapeutic group: Dermatologicals, antiseptics and disinfectants, aluminium agents.

ATC vet code: QD08AB

Micronised aluminium has long been used for the healing of wounds. It accelerates wound healing and has an antimicrobial effect due to the elimination of micro organisms from the tissues which it covers.

Aluspray allows a fine mist of aluminium to be produced permitting complete covering of lesions. The film thus obtained affords good protection against dirt and insects.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Liquid paraffin

Liquified petroleum gas

Stearic acid

6.2 Major incompatibilities

None known.

6.3 Shelf-life

Shelf life of the veterinary medicinal product as packaged for sale: 36 months.

6.4 Special precautions for storage

Do not store above 25°C.
Protect from direct sunlight.

6.5 Nature and composition of immediate packaging

This product is packed in an aluminium 210 ml pressurised can equipped with a valve.

6.6 Special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products

Do not pierce or burn even after use.

Any unused product or waste material should be disposed of in accordance with national requirements.

7 MARKETING AUTHORISATION HOLDER

Vetoquinol Ireland Limited
12 Northbrook Road
Ranelagh
Dublin 6.
Ireland

8 MARKETING AUTHORISATION NUMBER(S)

VPA10983/020/001

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 1st October 1989
Date of last renewal: 30th September 2009

10 DATE OF REVISION OF THE TEXT

August 2015