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VETOSPIRIN 1000 mg/g powder for use in drinking water/milk

Authorised

• Sodium salicylate

Product identification

Medicine name:

VETOSPIRIN 1000 mg/g powder for use in drinking water/milk Vetospirin 1000 mg/g Poudre pour administration dans le lait ou l'eau de boisson

Active substance:

Sodium salicylate

Target species:

Cattle (calf)

Pig

Route of administration:

In drinking water/milk use

Product details

Active substance and strength:

Sodium salicylate 1000.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Withdrawal period by route of administration: In drinking water/milk use:

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Cattle (calf)

- Meat and offal. no withdrawal period 0 days

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Pig

- Meat and offal. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

ON02BA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

- 1.0 kg multi-layered bag with an inner layer made of polyethylene
- 5.0 kg multi-layered bag with an inner layer made of polyethylene
- 10 multi-layer bags of 100 g with low density polyethylene inner layer packed in one cardboard box

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder: V.M.D.
Marketing authorisation date: 23/07/2019
Manufacturing sites for batch release: V.M.D.
Responsible authority: Ministry Of Health And Social Security
Authorisation number: V940/19/07/2001
Date of authorisation status change: 23/07/2019
Reference member state: Netherlands
Procedure number: NL/V/0250/001
Concerned member states: Belgium Estonia France Germany Hungary Latvia Lithuania Luxembourg Poland Portugal Romania Spain
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics

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