

Cryptisel 0.5 mg/ml oral solution for calves

Authorised

- Halofuginone lactate

Product identification

Medicine name:

Cryptisel 0.5 mg/ml oral solution for calves

Cryptisel vet 0,5 mg/ml Oral lösning

Active substance:

Halofuginone lactate

Target species:

Cattle (newborn calf)

Route of administration:

Oral use

Product details

Active substance and strength:

Halofuginone lactate

0.61 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:**Oral use:**

-

Cattle (newborn calf)

- Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51BX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Package description:

box containing 1 bottle of 1000 ml (containing 980 ml of solution)

box containing 1 bottle of 500 ml (containing 490 ml of solution)

box containing 1 bottle of 300 ml (containing 290 ml of solution)

box containing 1 bottle of 1000 ml (containing 980 ml of solution) with a 4 ml metering pump

box containing 1 bottle of 500 ml (containing 490 ml of solution) with a 4 ml metering pump

box containing 1 bottle of 300 ml (containing 290 ml of solution) with a 4 ml metering pump

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:

Industrial Veterinaria S.A.

Marketing authorisation date:

30/11/2020

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

aniMedica GmbH

aniMedica Herstellungs GmbH

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

59927

Date of authorisation status change:

30/11/2020

Reference member state:

Spain

Procedure number:

ES/V/0374/001

Concerned member states:

Austria Cyprus Czechia Denmark Estonia Germany Greece Hungary Ireland

Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania

Slovakia Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Package Leaflet

English (PDF)

Published on: 20/10/2025

[Download](#)

Summary of Product Characteristics

English (PDF)

Published on: 20/10/2025

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Combined File of all Documents

Labelling

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