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Kriptazen 0.5 mg/ml - Oral solution

Authorised

- Halofuginone

Product identification

Medicine name:
Kriptazen 0.5 mg/ml - Oral solution

Active substance:
Halofuginone

Target species:
Cattle (newborn calf)

Route of administration:
Oral use

Product details

Active substance and strength:
Halofuginone
0.50 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 13 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51AX

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:490 ml

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:980 ml

Packaging:Bottle (HDPE), Package_size:1 bottle + 1 metering pump (4 ml-12 ml), Content:490 ml

Packaging:Bottle (HDPE), Package_size:1 bottle + 1 metering pump (4 ml-12 ml), Content:980 ml

Packaging:Bottle (HDPE), Package_size:1 bottle + 1 metering pump (4 ml), Content:490 ml

Packaging:Bottle (HDPE), Package_size:1 bottle + 1 metering pump (4 ml), Content:980 ml

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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:
Virbac

Marketing authorisation date:
8/02/2019

Manufacturing sites for batch release:
Virbac

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
8/02/2019

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

Documents

Combined File of all Documents

English (PDF)

Published on: 14/03/2026

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