

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

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

Package description:

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

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, Content:980 ml

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

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 S.A.

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)

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