Cronyxin 50 mg/g Oral paste for horses

Authorised

- Flunixin meglumine

Product identification

Medicine name:
Cronyxin 50 mg/g Oral paste for horses

Active substance:
Flunixin meglumine

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Flunixin meglumine
83.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral paste

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 15 day
  - Milk. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AG90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Belgium

Package description:
(ID5) 396 gram(s): unspecified outer container with 12 Applicator (high-density polyethylene) each with 33 gram(s)
(ID4) 198 gram(s): unspecified outer container with 6 Applicator (high-density polyethylene) each with 33 gram(s)
(ID3) 99 gram(s): unspecified outer container with 3 Applicator (high-density polyethylene) each with 33 gram(s)
(ID2) 66 gram(s): unspecified outer container with 2 Applicator (high-density polyethylene) each with 33 gram(s)
(ID1) 33 gram(s): unspecified outer container with 1 Applicator (high-density polyethylene) with 33 gram(s)

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:
Bimeda Animal Health Limited

Marketing authorisation date:
22/01/2019

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V538240

Date of authorisation status change:
22/01/2019

Reference member state:
Germany

Procedure number:
DE/V/0178/001

Concerned member states:
Austria Belgium Denmark Estonia Finland France Ireland Italy Netherlands Poland Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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