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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:

Sweden

Available in:

Sweden

Package description:

(ID1) 33 gram(s): unspecified outer container with 1 Applicator (High Density PolyEthylene) with 33 gram(s)
(ID2) 66 gram(s): unspecified outer container with 2 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)
(ID4) 198 gram(s): unspecified outer container with 6 Applicator (High Density PolyEthylene) each with 33 gram(s)
(ID5) 396 gram(s): unspecified outer container with 12 Applicator (High Density PolyEthylene) each 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:
10/12/2018

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
56871

Date of authorisation status change:
10/12/2018

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

Combined File of all Documents

2402471-paren-20230821.pdf