

Cronyxin 50 mg/g Oral paste for horses

Authorised

- Flunixin meglumine

Product identification

Medicine name:

Cronyxin 50 mg/g Oral paste for horses

Cronyxin 50 mg/g pasta orale per equini

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:**

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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

Authorisation status:

Valid

Authorised in:

Italy

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:

1/08/2018

Manufacturing sites for batch release:

Bimeda Animal Health Limited

Responsible authority:

Ministry Of Health

Authorisation number:

105165

Date of authorisation status change:

1/08/2018

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

English (PDF)

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