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Flunipaste Vet. 50 mg/g Oral gel

Authorised

- Flunixin meglumine

Product identification

Medicine name:
Flunipaste Vet. 50 mg/g Oral gel

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 gel

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 15 day

Ej tillåtet för användning till lakterande djur som producerar mjölk för humankonsumtion

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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

Package description:
Available only in Swedish
Available only in Swedish
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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:
Biovet ApS

Marketing authorisation date:
12/12/2016

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
53882

Date of authorisation status change:
12/12/2016

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

Documents

Summary of Product Characteristics

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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.