Dexafort 2.67 mg/ml + 1.32 mg/ml suspension for injection for cattle, horses, dogs and cats

Authorised

- Dexamethasone sodium phosphate
- DEXAMETHASONE 21-PHENYLPROPIONATE

Product identification

Medicine name:
Dexafort 2.67 mg/ml + 1.32 mg/ml suspension for injection for cattle, horses, dogs and cats

Active substance:
Dexamethasone sodium phosphate
DEXAMETHASONE 21-PHENYLPROPIONATE

Target species:
Cattle
Horse
Cat
Dog

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
1.32 milligram(s) / 1.00 millilitre(s)

DEXAMETHASONE 21-PHENYLPROPIONATE
2.67 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 53 day
  - Milk. 6 day Мляко: 5.5 дни (11 издоявания)
- **Horse**
  - Meat and offal. 47 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
20/02/2008

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1956

Date of authorisation status change:
26/02/2013

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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