Dexafort, 3 mg/ml süstesuspensioon veistele, hobustele, koertele ja kassidele

Not authorised

- Dexamethasone sodium phosphate
- DEXAMETHASONE 21-PHENYLPROPIONATE

Product identification

Medicine name:
Dexafort, 3 mg/ml süstesuspensioon veistele, hobustele, koertele ja kassidele

Active substance:
Dexamethasone sodium phosphate
DEXAMETHASONE 21-PHENYLPROPIONATE

Target species:
Horse
Cattle
Dog
Cat

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:

- **Horse**
  - Meat and offal. 47 day
  - Milk. no withdrawal period

Mitte kasutada märadel, kelle piima kavatsetakse tarvitada inimtoiduks.

- **Cattle**
  - Meat and offal. 53 day
  - Milk. 6 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:
Surrendered

Authorised in:
Estonia

Package description:
Available only in Estonian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
5/02/2004

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
State Agency Of Medicines

Authorisation number:
1186

Date of authorisation status change:
25/02/2024

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.