DEXAFORT (1,32+2,67)MG/ML ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Not authorised

- DEXAMETHASONE 21-PHENYLPROPIONATE
- Dexamethasone sodium phosphate

Product identification

Medicine name:
DEXAFORT (1,32+2,67)MG/ML ΕΝΕΣΙΜΟ ΕΝΑΙΩΡΗΜΑ

Active substance:
DEXAMETHASONE 21-PHENYLPROPIONATE
Dexamethasone sodium phosphate

Target species:
Horse
Cattle
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

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

Dexamethasone sodium phosphate
1.32 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
- **Cattle**
  - Meat and offal. 53 day
  - Milk. 7 day
- **Dog**
  - Not applicable. no withdrawal period
- **Cat**
  - Not applicable. no withdrawal period

Subcutaneous use:

- **Dog**
  - Not applicable. no withdrawal period
- **Cat**
  - Not applicable. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

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

Authorisation status:
Surrendered

Authorised in:
Greece

Package description:
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
2/10/2000

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

Responsible authority:
National Organization For Medicines

Authorisation number:
37940/3-10-2000/K-0129201

Date of authorisation status change:
22/05/2024

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