Dexafort Suspensie voor injectie

Authorised

- Dexamethasone sodium phosphate
- DEXAMETHASONE 21-PHENYLPROPIONATE

Product identification

Medicine name:
Dexafort Suspensie voor injectie

Active substance:
Dexamethasone sodium phosphate
DEXAMETHASONE 21-PHENYLPROPIONATE

Target species:
Dog
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
1.47 milligram(s) / 1.00 millilitre(s)
DEXAMETHASONE 21-PHENYLPROPIONATE

2.70 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 63 day
  - Milk. no withdrawal period 14 milkings

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Dexafort 50 ml Vial (glass) Suspension for injection

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:
1/05/1972

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V059437

Date of authorisation status change:
5/12/2023

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.