

DEXAFORT

Authorised

- DEXAMETHASONE 21-PHENYLPROPIONATE
- DEXAMETHASONE DISODIUM PHOSPHATE

Product identification

Medicine name:

DEXAFORT

Active substance:

DEXAMETHASONE 21-PHENYLPROPIONATE

DEXAMETHASONE DISODIUM PHOSPHATE

Target species:

Horse

Cattle

Dog

Route of administration:

Intramuscular use

Product details

Active substance and strength:

DEXAMETHASONE 21-PHENYLPROPIONATE

1.32 milligram(s) / 1.00 millilitre(s)

DEXAMETHASONE DISODIUM PHOSPHATE

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. 48 day

Use in mares producing milk for human consumption is not permitted

-

Cattle

- Meat and offal. 48 day

- Milk. 132 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

30/03/2010

Manufacturing sites for batch release:

Vet Pharma Friesoythe GmbH

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

160115

Date of authorisation status change:

12/08/2025

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

Documents

Combined File of all Documents

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