Rapidexon 2 mg/ml solution for injection

• Dexamethasone sodium phosphate

Product identification

Medicine name:

Rapidexon 2 mg/ml solution for injection Rapidexon 2 mg/ml injekcinis tirpalas

Active substance:

Dexamethasone sodium phosphate

Target species:

Horse Cattle

Pig

Dog

Cat

Route of administration:

Intraarticular use Intramuscular use Intravenous use Intrabursal use

Product details

Active substance and strength:

Dexamethasone sodium phosphate 2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration: Intraarticular use:

Horse

•

•

- Meat and offal. 8 day

Intramuscular use:

Cattle

- Milk. 72 hour
- Meat and offal. 8 day

Pig

- Meat and offal. 2 day

Horse

- Meat and offal. 8 day

•

Dog

•

Cat

•

•

Intravenous use:

Horse

- Meat and offal. 8 day

Intrabursal use:

Horse

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

50 ml vial glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium cap

25 ml vial (filled in 30 ml vial) glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium cap

100 ml vial glass type I uncoloured with bromobutyl rubber stopper type I secured with aluminium

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Eurovet Animal Health B.V.

Marketing authorisation date:

25/03/2008

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

State Food And Veterinary Service

Authorisation number: LT/2/08/1787/001-003

Date of authorisation status change:

30/07/2011

Reference member state:

Netherlands

Procedure number: NL/V/0284/001

Concerned member states:

Austria Belgium Czechia Denmark Finland France Greece Hungary Ireland Lithuania Poland Portugal Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

RV1787.pdf

Rapidexon 2 mg.pdf

Source URL: *https://medicines.health.europa.eu/veterinary/60000037289*