

Rapidexon 2 mg/ml Solution injectable

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

Rapidexon 2 mg/ml Solution injectable

Rapidexon 2 mg/ml Injektionslösung

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Pig

Horse

Dog

Cat

Route of administration:

Intraarterial use

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate

2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intraarterial use:

-

Cattle

- Meat and offal. 8 day

- Milk. 72 hour

-

Pig

- Meat and offal. 2 day

-

Horse

- Meat and offal. 8 day

Intramuscular use:

-

Cattle

- Meat and offal. 8 day

- Milk. 72 hour

-

Pig

- Meat and offal. 2 day

-

Horse

- Meat and offal. 8 day

Intravenous use:

•

Cattle

- Meat and offal. 8 day

- Milk. 72 hour

•

Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

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

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

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

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:

Dechra Regulatory B.V.

Marketing authorisation date:

4/02/1998

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:

V 914/98/02/0586

Date of authorisation status change:

24/03/2008

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.