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

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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

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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.