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Rapidexon 2 mg/ml oplossing voor injectie

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
Rapidexon 2 mg/ml oplossing voor injectie

Active substance:
Dexamethasone sodium phosphate

Target species:
Cattle
Pig
Horse
Dog
Cat

Route of administration:
Intramuscular 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:

Intramuscular use:

- **Cattle**
  - Milk. 72 hour
  - Meat and offal. 8 day
- **Pig**
  - Meat and offal. 2 day
- **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:
Netherlands

Available in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch
Available only in Dutch

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:
6/08/2009

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 105653

Date of authorisation status change:
19/06/2014

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

Documents

Combined File of all Documents

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