FATROCORTIN 1mg/ml - ενέσιμο διάλυμα για βοοειδή, χοίρους, άλογα, σκύλους και γάτους

Authorised

- Dexamethasone

Product identification

Medicine name:
FATROCORTIN 1mg/ml - ενέσιμο διάλυμα για βοοειδή, χοίρους, άλογα, σκύλους και γάτους

Active substance:
Dexamethasone

Target species:
Cattle
Dog
Horse
Cat
Pig

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:

Dexamethasone
1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 16 day
  - Milk. 72 hour
- **Dog**
- **Horse**
  - Meat and offal. 24 day
- **Cat**
- **Pig**
  - Meat and offal. 3 day

Intravenous use:

- **Cattle**
  - Meat and offal. 16 day
  - Milk. 72 hour
- **Pig**
  - Meat and offal. 3 day
- **Horse**
  - Meat and offal. 24 day
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Cyprus

Available in:
Cyprus

Package description:
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
21/11/1985

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Agriculture Rural Development And Environment

Authorisation number:
10310

Date of authorisation status change:
6/08/2012

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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