

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

-

Horse

- Meat and offal. 24 day

-

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product 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:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

10310

Date of authorisation status change:

21/11/1985

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