ADECON

Authorised

- TOCOPHERYL ACETATE
- RETINOL ACETATE
- Colecalciferol

Product identification

Medicine name:
ADECON

Active substance:
TOCOPHERYL ACETATE

RETINOL ACETATE

Colecalciferol

Target species:
Cattle
Goat
Sheep
Pig
Horse (food producing)

Route of administration:
Oral use
Intramuscular use

Product details

Active substance and strength:

TOCOPHERYL ACETATE
100.00 milligram(s) / 1.00 millilitre(s)

RETINOL ACETATE
100000.00 international unit(s) / 1.00 millilitre(s)

Colecalciferol
25000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Milk. 120 hour

- **Cattle**
  - Meat and offal. 243 day

- **Goat**
  - Milk. 120 hour

- **Goat**
  - Meat and offal. 215 day

- **Sheep**
  - Milk. 120 hour

- **Sheep**
  - Meat and offal. 215 day

- **Pig**

- Meat and offal. 215 day

- **Horse (food producing)**
  - Milk. 120 hour

- **Horse (food producing)**
  - Meat and offal. 243 day

Intramuscular use:

- **Cattle**
  - Milk. 120 hour

- **Cattle**
  - Meat and offal. 243 day

- **Goat**
  - Milk. 120 hour

- **Goat**
  - Meat and offal. 215 day

- **Sheep**
  - Milk. 120 hour

- **Sheep**
  - Meat and offal. 215 day

- **Pig**
  - Meat and offal. 215 day

- **Horse (food producing)**
  - Milk. 120 hour

- **Horse (food producing)**
  - Meat and offal. 243 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
16/02/1974

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

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
19/11/2009

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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