

# ADECON

Authorised

- TOCOPHERYL ACETATE
- RETINOL ACETATE
- Colecalciferol

## Product identification

**Medicine name:**

ADECON

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**Active substance:**

TOCOPHERYL ACETATE

RETINOL ACETATE

Colecalciferol

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**Target species:**

Cattle

Goat

Sheep

Pig

Horse (food producing)

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**Route of administration:**

Oral use

Intramuscular use

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## 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)

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### Pharmaceutical form:

Solution for injection

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### Withdrawal period by route of administration:

#### Oral use:

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#### Cattle

- Milk. 120 hour

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#### Cattle

- Meat and offal. 243 day

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#### Goat

- Milk. 120 hour

- 

#### Goat

- Meat and offal. 215 day

- 

#### Sheep

- Milk. 120 hour

- 

#### Sheep

- Meat and offal. 215 day

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**Pig**

- Meat and offal. 215 day

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA11JA

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Package description:**

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

16/02/1974

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

19/11/2009

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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