

# ADECON INJECTION Ενέσιμο διάλυμα για βοοειδή, άλογα, χοίρους, πρόβατα και αίγες

Authorised

- alpha-Tocopheryl acetate concentrate (powder form)
- RETINOL ACETATE
- Colecalciferol

## Product identification

### **Medicine name:**

ADECON INJECTION Ενέσιμο διάλυμα για βοοειδή, άλογα, χοίρους, πρόβατα και αίγες

### **Active substance:**

alpha-Tocopheryl acetate concentrate (powder form)

RETINOL ACETATE

Colecalciferol

### **Target species:**

Cattle (calf)

Goat

Sheep

Horse

Pig

### **Route of administration:**

Intramuscular use

Oral use

## Product details

### **Active substance and strength:**

alpha-Tocopheryl acetate concentrate (powder form)

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:**

#### **Intramuscular use:**

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#### **Cattle (calf)**

- Milk. 0 day

- Meat and offal. 0 day

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#### **Goat**

- Meat and offal. 0 day

- Milk. 0 day

•

#### **Sheep**

- Meat and offal. 0 day

- Milk. 0 day

•

#### **Horse**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Pig**

- Meat and offal. 0 day

**Oral use:**

- 

**Cattle (calf)**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Goat**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Sheep**

- Meat and offal. 0 day

- Milk. 0 day

- 

**Horse**

- Meat and offal. 0 day

- Milk. 0 day

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**Pig**

- Meat and offal. 0 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:**

Cyprus

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**Available in:**

Cyprus

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**Package description:**

Available only in Greek

Available only in Greek

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (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:**

20/01/1988

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

11599

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**Date of authorisation status change:**

20/01/1988

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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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