

# Clamoxyl Vet. injektionsvæske, suspension 150 mg/ml

Authorised

- Amoxicillin trihydrate
- Amoxicillin trihydrate

## Product identification

### **Medicine name:**

Clamoxyl Vet. injektionsvæske, suspension 150 mg/ml

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### **Active substance:**

Amoxicillin trihydrate

Amoxicillin trihydrate

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### **Target species:**

Cattle

Pig

Cat

Sheep

Dog

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### **Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

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

Amoxicillin trihydrate

183.00 milligram(s) / 1.00 millilitre(s)

Amoxicillin trihydrate

183.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Suspension for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Cattle**

- Milk. 3 day
- Milk. 3 day
- Meat and offal. 30 day
- Meat and offal. 30 day

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#### **Pig**

- Meat and offal. 30 day
- Meat and offal. 30 day

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#### **Sheep**

- Meat and offal. 30 day
- Meat and offal. 30 day

#### **Subcutaneous use:**

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#### **Pig**

- Meat and offal. 30 day

- Meat and offal. 30 day

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**Sheep**

- Meat and offal. 30 day

- Meat and offal. 30 day

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**Cattle**

- Milk. 3 day

- Milk. 3 day

- Meat and offal. 30 day

- Meat and offal. 30 day

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

QJ01CA04

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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:**

Denmark

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**Available in:**

Denmark

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**Package description:**

Available only in [Danish](#)

Available only in [Danish](#)

Available only in [Danish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Zoetis Animal Health ApS

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**Marketing authorisation date:**

13/04/1994

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**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

15053

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**Date of authorisation status change:**

13/04/1994

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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

Summary of Product Characteristics

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