

# Vetrimoxin L.A. 150 mg/ml Suspension for Injection for Cattle and Pigs

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Vetrimoxin L.A. 150 mg/ml Suspension for Injection for Cattle and Pigs

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**Active substance:**

Amoxicillin trihydrate

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Amoxicillin trihydrate

172.40 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
- Meat and offal. 18 day

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**Pig**

- Meat and offal. 16 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:**

United Kingdom (Northern Ireland)

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**Package description:**

(ID1) 1200 millilitre(s): unspecified outer container with 12 Vial (Plastic) each with 100 millilitre(s)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Plastic) with 100 millilitre(s)

(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Plastic) with 250 millilitre(s)

(ID4) 3000 millilitre(s): unspecified outer container with 12 Vial (Plastic) each with 250 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

19/12/2012

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 14966/3064

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**Date of authorisation status change:**

27/06/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0153/001

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**Concerned member states:**

Austria Denmark Finland Iceland Ireland Netherlands Sweden

United Kingdom (Northern Ireland)

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