

# INTRAMICINE

Authorised

- Benzylpenicillin procaine monohydrate
- Dihydrostreptomycin sulfate

## Product identification

**Medicine name:**

INTRAMICINE

**Active substance:**

Benzylpenicillin procaine monohydrate

Dihydrostreptomycin sulfate

**Target species:**

Cattle

Pig

Cat

Sheep

Goat

Dog

**Route of administration:**

Intramuscular use

Subcutaneous use

## Product details

**Active substance and strength:**

Benzylopenicillin procaine monohydrate

200.00 milligram(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate

250.40 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**

- Meat and offal. 30 day
- Milk. 7 day

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**Pig**

- Meat and offal. 30 day

- 

**Sheep**

- Meat and offal. 30 day
- Milk. 6 day

- 

**Goat**

- Meat and offal. 30 day
- Milk. 7 day

**Subcutaneous use:**

- 

**Cattle**

- Meat and offal. 30 day
- Milk. 7 day

- 

**Pig**

- Meat and offal. 30 day

- 

#### **Sheep**

- Meat and offal. 30 day

- Milk. 6 day

- 

#### **Goat**

- Meat and offal. 30 day

- Milk. 7 day

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

QJ01RA01

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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:**

France

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#### **Package description:**

Available only in French

Available only in French

Available only in French

Available only in French

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## **Additional information**

#### **Entitlement type:**

Marketing Authorisation

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#### **Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

30/09/1988

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**Manufacturing sites for batch release:**

CEVA SANTE ANIMALE - LIBOURNE

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/7100510 2/1988

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**Date of authorisation status change:**

30/09/2013

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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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Package Leaflet and Labelling

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