**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000035483

# **AMPICLOX**

Authorised

- Ampicillin sodium
- Cloxacillin sodium monohydrate

## Product identification

#### **Medicine name:**

**AMPICLOX** 

#### **Active substance:**

Ampicillin sodium

Cloxacillin sodium monohydrate

## **Target species:**

Cattle (cow)

#### **Route of administration:**

Intramammary use

## **Product details**

## **Active substance and strength:**

Ampicillin sodium 79.78 milligram(s) / 1.00 Syringe

Cloxacillin sodium monohydrate

218.35 milligram(s) / 1.00 Syringe

#### **Pharmaceutical form:**

# Withdrawal period by route of administration: Intramammary use:

Cattle (cow)

- Meat and offal. 7 day
- Milk. 48 hour

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

QJ51RC26

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

France

## Package description:

Available only in <u>French</u> Available only in French

## Additional information

# **Entitlement type:**

Marketing Authorisation

## Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

# Marketing authorisation holder:

Zoetis France

## Marketing authorisation date:

6/05/1988

### Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

## **Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

#### **Authorisation number:**

FR/V/8027594 3/1988

## Date of authorisation status change:

6/05/2013

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## **Documents**

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.