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:

Intramammary suspension

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 French
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

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Package Leaflet and Labelling

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