AMPICLOX LC, intramaminė suspensija galvijams (karvėms laktacijos metu)

Authorised

- Ampicillin sodium
- Cloxacillin sodium

Product identification

Medicine name:
AMPICLOX LC, intramaminė suspensija galvijams (karvėms laktacijos metu)

Active substance:
Ampicillin sodium

Cloxacillin sodium

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Ampicillin sodium
75.00 milligram(s) / 1.00 Syringe

Cloxacillin sodium
200.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle**
  - Meat and offal. 7 day
  - Milk. 60 hour

When milking cows twice a day, milk can only be used for human consumption after 60 hours. (i.e. 5 milkings) after the last treatment. The milk may be used for human consumption only after the same period after the last treatment (for example, by milking three times a day and after taking the medicine twice a day, milk can only be used for human consumption from the 8th milking).

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CR50

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
22/02/2000

Manufacturing sites for batch release:
Haupt Pharma Latina S.r.l.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/00/1073/001-002

Date of authorisation status change:
27/05/2024

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

Documents

Combined File of all Documents

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