SECLARIS DC 250 MG INTRAMAMMARY SUSPENSION FOR DRY COWS

Not authorised

- Cefalonium dihydrate

Product identification

Medicine name:
SECLARIS DC 250 MG INTRAMAMMARY SUSPENSION FOR DRY COWS
Seclaris DC, 250 mg intramaminė suspensija užtrūkusioms karvėms

Active substance:
Cefalonium dihydrate

Target species:
Cattle (cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalonium dihydrate
269.60 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (cow)**
  - Milk. 96 hour 96 hours after calving if the dry period is longer than 54 days
  - Meat and offal. 21 day
  - Milk. 58 day

58 days following the treatment if the dry period is less than or equal to 54 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB90

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

Authorisation status:
Surrendered

Authorised in:
Lithuania

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
25/10/2017

Manufacturing sites for batch release:
Lohmann Pharma Herstellung GmbH

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/17/2426/001-002

Date of authorisation status change:
25/10/2017

Reference member state:
France

Procedure number:
FR/V/0399/001

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

Documents

RV2426.pdf