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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, intramammális szuszpenzió szárazonálló tehenek részére A.U.V.

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:

Hungary

Package description:

72 intramammary syringes and 72 cleaning towels

20 intramammary syringes and 20 cleaning towels

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-Phylaxia Zrt.
Marketing authorisation date: 22/11/2017
Manufacturing sites for batch release: Lohmann Pharma Herstellung GmbH
Responsible authority: Directorate Of Veterinary Medicinal Products
Authorisation number: 3909/X/2017 NÉBIH ÁTI
Date of authorisation status change: 25/04/2024
Reference member state: France
Procedure number: FR/V/0399/001

To consult adverse reactions on veterinary medicinal products please go to $\underline{www.adrreports.eu/vet}$