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ORBENIN DC 500 mg intramamarna suspenzija za krave v presušitvi

Authorised

- Cloxacillin

Product identification

Medicine name:
ORBENIN DC 500 mg intramamarna suspenzija za krave v presušitvi

Active substance:
Cloxacillin

Target species:
Cattle (cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cloxacillin
500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (cow)**
  - Meat. 0 day Meso: 0 dni
  - Milk. 4 day

Mleko: 4 dni po telitvi. Če se krava predčasno oteli ali povrže, oziroma če ji zdravilo aplicirate v obdobju, krajšem od 28 dni do datuma telitve, je mleko primerno za prehrano 28. dan plus 96 ur po vnosu zdravila

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Available in:

Slovenia

Package description:

Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
1/06/2001

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0244/001

Date of authorisation status change:
1/06/2001

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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