Orbenin extra dry cow 600 mg suspensie voor intramammair gebruik voor runderen

Authorised

- Cloxacillin hemibenzathine

Product identification

Medicine name:
Orbenin extra dry cow 600 mg suspensie voor intramammair gebruik voor runderen

Active substance:
Cloxacillin hemibenzathine

Target species:
Cattle

Route of administration:
Intramammary use

Product details

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

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Milk. 44 day

44 days after last treatment, after calving within 42 days after treatment

  - Milk. 48 hour 48 hours after calving after more than 42 days after treatment
  - Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

Available only in Dutch

Available only in Dutch

Available only in Dutch

Additional information

Entitlement type:

Marketing Authorisation

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

Marketing authorisation holder:
Zoetis B.V.

Marketing authorisation date:
9/11/1989

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 6901

Date of authorisation status change:
22/11/2016

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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