DRYCLOX-N

Authorised

- NEOMYCIN SULFATE
- Cloxacillin hemibenzathine

Product identification

Medicine name:
DRYCLOX-N

Active substance:
NEOMYCIN SULFATE

Cloxacillin hemibenzathine

Target species:
Cattle (dairy cow at drying-off)

Route of administration:
Intramammary use

Product details

Active substance and strength:
NEOMYCIN SULFATE
250.00 milligram(s) / 1.00 Syringe

Cloxacillin hemibenzathine
500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dairy cow at drying-off)**
  - Meat and offal. 70 day
  - Milk. 45 day 45 дни при сухостоен период по-къс от 6 седмици
  - Milk. 3 day

3 дни при сухостоен период от 6 - 8 седмици, 2 дни при сухостоен период по-дълъг от 8 седмици

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC26

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

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:
Kepro B.V.

Marketing authorisation date:
9/02/2012

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1708

Date of authorisation status change:
19/12/2016

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

Documents

Summary of Product Characteristics

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