Multishield DC Intramammary Suspension for Cows

Authorised

- NEOMYCIN SULFATE
- Penethamate hydriodide
- Benzylpenicillin procaine

Product identification

Medicine name:
Multishield DC Intramammary Suspension for Cows
Multishield DC Suspension zur intramammären Anwendung bei Rindern

Active substance:
NEOMYCIN SULFATE

Penethamate hydriodide

Benzylpenicillin procaine

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:

NEOMYCIN SULFATE
100.00 milligram(s) / 1.00 Syringe

Penethamate hydriodide
100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine
400.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 96 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RC22

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

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Authorisation status:
Valid

Authorised in:
Germany

Package description:
Low density Polyethylene intramammary syringe, containing 4.5g intramammary suspension.Syringes packed in cartons of 24 syringes.
Low density Polyethylene intramammary syringe, containing 4.5g intramammary suspension.Syringes packed in buckets of 120 syringes.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
1/03/2013

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401543.00.00

Date of authorisation status change:
15/08/2018

Reference member state:
Ireland

Procedure number:
IE/V/0277/001

Concerned member states:
Belgium Czechia France Germany Hungary Italy Netherlands Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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