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KEFAMAST DRY COW, intramaminė suspensija karvėms

Not authorised

- Dihydrostreptomycin
- Cefalexin

Product identification

Medicine name:
KEFAMAST DRY COW, intramaminė suspensija karvėms

Active substance:
Dihydrostreptomycin

Cefalexin

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

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

Cefalexin

500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 28 day
  - Milk. 3 day

If the cow calves less than 40 days later. after treatment for milk - 42.5 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RD01

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

Authorisation status:
Surrendered

Authorised in:
Lithuania

Package description:
Polietileniniai švirkštai po 10 ml (9 g) intramaminės suspensijos, kibirėliuose po 120 vnt.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
18/08/1998

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/98/0723/001

Date of authorisation status change:
28/04/2009

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

Documents

RV0723.pdf