FATROXIMIN dry cow, 100 mg/5 ml intramammary ointment for cows and buffalo cows

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN dry cow, 100 mg/5 ml intramammary ointment for cows and buffalo cows

Active substance:
Rifaximin

Target species:
Buffalo
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Rifaximin
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary ointment

Withdrawal period by route of administration:
Intramammary use:

- **Buffalo**
  - Milk. 0 hour

Μηδέν ώρες (μηδέν αρμέγματα) μετά τον τοκετό, εάν η περίοδος ξηρασίας είναι ίση ή μεγαλύτερη από 49 ημέρες, 49 ημέρες μετά τη θεραπεία εάν η περίοδος ξηρασίας είναι μικρότερη από 49 ημέρες.

  - Meat and offal. 0 day

Να μην χρησιμοποιούνται οι μαστοί των ζώων που υποβάλλονται σε θεραπεία για ανθρώπινη κατανάλωση.

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51XX01

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

Authorisation status:
Valid

Authorised in:
Cyprus

Available in:
Cyprus

Package description:
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
30/04/2004

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00078V

Date of authorisation status change:
12/05/2014

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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