Bimastat Oral Suspension

Authorised

- NEOMYCIN SULFATE
- Sulfadiazine

Product identification

Medicine name:
Bimastat Oral Suspension

Active substance:
NEOMYCIN SULFATE

Sulfadiazine

Target species:
Cattle

Route of administration:
Oral use

Product details

Active substance and strength:
NEOMYCIN SULFATE
28.99 milligram(s) / 1.00 millilitre(s)

Sulfadiazine
150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01RA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

1 litre white high density polythylene bottles, sealed with a tamper-evident cap containing a pink oralsuspension.
400 ml white high density polythylene bottles, sealed with a tamper-evident cap containing a pink oralsuspension.
250 ml white high density polythylene bottles, sealed with a tamper-evident cap containing a pink oralsuspension.A 50 ml dosing syringe is supplied with the 250 ml presentation only.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bimeda Animal Health Limited

Marketing authorisation date:
1/10/1988

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22033/001/001

Date of authorisation status change:
1/10/1988

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

Documents

Summary of Product Characteristics