Norodine Equine Oral Paste

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:
Norodine Equine Oral Paste

Active substance:
Trimethoprim

Sulfadiazine

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Trimethoprim
5.80 gram(s) / 100.00 gram(s)

Sulfadiazine
28.83 gram(s) / 100.00 gram(s)

Pharmaceutical form:

Oral paste

Withdrawal period by route of administration:

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Norodine Equine Paste is presented in a white high density polyethylene oral syringe with a high density polyethylene plungercalibrated in divisions, each equivalent to 50kg of bodyweight and a white high density polyethylene cap (push-fit). Eachsyringe contains enough for one daily dose for a 500kg horse.

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:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
1/10/1988

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/019/001

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