

Norbonex 5 mg/ml Pour-on Solution for Beef and Dairy Cattle

Not authorised

- Eprinomectin

Product identification

Medicine name:

Norbonex 5 mg/ml Pour-on Solution for Beef and Dairy Cattle

Active substance:

Eprinomectin

Target species:

Cattle (for meat production)

Cattle (dairy cow)

Route of administration:

Pour-on use

Product details

Active substance and strength:

Eprinomectin

5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Pour-on solution

Withdrawal period by route of administration:**Pour-on use:**

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Cattle (for meat production)

- Meat and offal. 10 day

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Cattle (dairy cow)

- Meat and offal. 10 day

- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

United Kingdom (Northern Ireland)

Package description:

(ID5) 1 litre(s): unspecified outer container with 1 Barrel (Polyethylen hoher Dichte) with 1 litre(s)

(ID4) 5 litre(s): unspecified outer container with 1 Barrel (Polyethylen hoher Dichte) with 5 litre(s)

(ID3) 2.5 litre(s): unspecified outer container with 1 Barrel (Polyethylen hoher Dichte) with 2.5 litre(s)

(ID2) 1 litre(s): unspecified outer container with 1 Container (Polyethylen hoher Dichte) with 1 litre(s), closed with Schnappdeckel (Polyethylen hoher Dichte)

(ID1) 250 millilitre(s): unspecified outer container with 1 Container (Polyethylen hoher Dichte) with 250 millilitre(s), closed with Schnappdeckel (Polyethylen hoher Dichte)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories Limited

Marketing authorisation date:

11/06/2014

Manufacturing sites for batch release:

Norbrook Laboratories (Ireland) Limited

Norbrook Laboratories Limited

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 02000/4342

Date of authorisation status change:

11/07/2022

Reference member state:

Germany

Procedure number:

DE/V/0327/001

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