

Enovex 10 mg/ml Solution for Injection for cattle, sheep and pigs

Authorised

- Ivermectin

Product identification

Medicine name:

Enovex 10 mg/ml Solution for Injection for cattle, sheep and pigs

Active substance:

Ivermectin

Target species:

Cattle
Sheep
Pig

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Cattle

- Meat and offal. 49 day

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Sheep

- Meat and offal. 42 day

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Pig

- Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

The product will be supplied in 1L volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 500ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 250ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 100ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

The product will be supplied in 50ml volume, presented in high density polyethylene vial with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

29/10/1999

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22664/055/001

Date of authorisation status change:

29/10/1999

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

Documents

Summary of Product Characteristics