Bovimectin 10 mg/ml solution for injection for cattle

Not authorised

- Ivermectin

Product identification

Medicine name:
Bovimectin 10 mg/ml solution for injection for cattle
Bovimectin 10 mg/ml solution for injection for cattle

Active substance:
Ivermectin

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
1.00 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 49 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
Multidose high density polyethylene bottle of 50 ml sealed with bromobutyl seal and plain aluminiumoverseal. Cardboard box with 1 x 50 ml bottle.
Multidose high density polyethylene bottle of 250 ml sealed with bromobutyl seal and plain aluminiumoverseal. Cardboard box with 1 x 250 ml bottle.
Multidose high density polyethylene bottle of 500 ml sealed with bromobutyl seal and plain aluminiumoverseal. Cardboard box with 1 x 500 ml bottle.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
5/06/2001

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22033/051/001

Date of authorisation status change:
31/01/2025

Reference member state:
Spain

Procedure number:
ES/V/0458/001

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

Documents

Summary of Product Characteristics