Intermectin Injection 10 mg/ml solution for injection for cattle, sheep and pigs

Authorised

- Ivermectin

Product identification

Medicine name:
Intermectin Injection 10 mg/ml solution for injection for cattle, sheep and pigs
INTERMECTIN INJECTION 10 MG/ML SOLUTION INJECTABLE POUR BOVINS OVINS ET PORCINS

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:

- **Cattle**
  - Meat and offal. 49 day
- **Sheep**
  - Meat and offal. 63 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Clear glass bottles (type II) closed with a bromobutyl rubber stopper and sealed with an aluminium cap packed into an outer cardboard box
Clear glass bottles (type II) closed with a bromobutyl rubber stopper and sealed with an aluminium flip-off cap with polypropylene cover packed into an outer cardboard box

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Interchemie Werken De Adelaar B.V.

Marketing authorisation date:
20/12/2024

Manufacturing sites for batch release:
Interchemie Werken De Adelaar Eesti AS

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5189648 1/2024

Date of authorisation status change:
20/12/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0431/001

Concerned member states:
Austria Croatia Cyprus Czechia France Germany Greece Hungary Italy Latvia Lithuania Malta Portugal Slovakia Slovenia Spain

Generic of:
600000065934

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

Documents

Combined File of all Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.