Source URL: https://medicines.health.europa.eu/veterinary/en/60000014300

IVOMEC PLUS, soluție injectabilă pentru bovine

Authorised

- Clorsulon
- Ivermectin

Product identification

Medicine name:

IVOMEC PLUS, soluție injectabilă pentru bovine

Active substance:

Clorsulon

Ivermectin

Target species:

Cattle

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Clorsulon

100.00 milligram(s) / 1.00 millilitre(s)

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. 66 day

The product should not be administered to animals during the period of milk production or 28 days before calving if the milk is intended for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

4/08/2003

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France SCS

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

120072

Date of authorisation status change:

22/06/2025

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

Documents

Combined File of all Documents

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