

LEVAVERMIN 50 mg/ml

Authorised

- Levamisole hydrochloride

Product identification

Medicine name:

LEVAVERMIN 50 mg/ml

Active substance:

Levamisole hydrochloride

Target species:

Cattle

Sheep

Pig

Chicken (broiler)

Route of administration:

Oral use

Product details

Active substance and strength:

Levamisole hydrochloride

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

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

-

Cattle

- Meat and offal. 3 day

Nu este autorizata utilizarea la animalele care produc lapte pentru consum uman

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Sheep

- Meat and offal. 3 day

Nu este autorizata utilizarea la animalele care produc lapte pentru consum uman

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Pig

- Meat and offal. 3 day

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Chicken (broiler)

- Meat and offal. 3 day

Nu este autorizata utilizarea la gainile care produc oua pentru consum uman.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in [Romanian](#)
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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Crida Pharm S.R.L.

Marketing authorisation date:

7/09/2015

Manufacturing sites for batch release:

Crida Pharm S.R.L.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

200097

Date of authorisation status change:

24/10/2022

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

Documents

Summary of Product Characteristics

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