

Ecomectin, 10.0mg/ml, Injekční roztok

Authorised

- Ivermectin

Product identification

Medicine name:

Ecomectin, 10.0mg/ml, Injekční roztok

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

- Milk. no withdrawal period

Nepoužívat u skotu méně než 60 dní před otelením.,

- Meat and offal. 49 day

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Sheep

- Milk. no withdrawal period

Nepoužívat u zvířat, jejichž mléko je určeno pro lidskou spotřebu.,

- Meat and offal. 42 day

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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:

Czechia

Package description:

Available only in [Czech](#)

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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:

Eco Animal Health Europe Limited

Marketing authorisation date:

29/12/2000

Manufacturing sites for batch release:

Eco Animal Health Limited

Divasa Farmavic S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/092/00-C

Date of authorisation status change:

29/12/2000

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.

Package Leaflet

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

Labelling

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