

Ivomec Super 10,0 mg/ml + 100,0 mg/ml injekčný roztok

Authorised

- Clorsulon
- Ivermectin

Product identification

Medicine name:

Ivomec Super 10,0 mg/ml + 100,0 mg/ml injekčný roztok

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:

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Cattle

- Meat and offal. 66 day

- Milk. 0 day

Do not use in animals producing milk 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:

Slovakia

Package description:

Available only in [Slovak](#)

Available only in [Slovak](#)

Available only in [Slovak](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

13/06/1994

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/190/90-S

Date of authorisation status change:

13/06/1994

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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