

BIOMECTIN 10 mg/ml injekčný roztok

Authorised

- Ivermectin

Product identification

Medicine name:

BIOMECTIN 10 mg/ml injekčný 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

- Meat and offal. 28 day

Do not use in animals which milk is produced for human consumption.

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Sheep

- Meat and offal. 28 day

Do not use in animals which milk is produced for human consumption.

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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:

Slovakia

Available in:

Slovakia

Package description:

Available only in Slovak

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:

Vetoquinol s.r.o.

Marketing authorisation date:

30/05/1997

Manufacturing sites for batch release:

Vetoquinol Biowet Sp. z o.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/0074/97-S

Date of authorisation status change:

30/05/1997

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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