

Santiola 50 mg/ml solution for injection for cattle and sheep

Not
authorised

- Closantel sodium dihydrate

Product identification

Medicine name:

Santiola 50 mg/ml solution for injection for cattle and sheep
Santiola 50 mg/ml injekčný roztok pre hovädzí dobytok a ovce

Active substance:

Closantel sodium dihydrate

Target species:

Cattle
Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Closantel sodium dihydrate
54.38 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. 77 day

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Sheep

- Meat and offal. 107 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Slovakia

Package description:

Container or pack size: 4 type I amber glass vials. Closure: bromobutyl rubber.

Contents of each vial: 250 ml solution.

Container or pack size: 1 type I amber glass vial. Closure: bromobutyl rubber.

Contents of each vial: 250 ml solution.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

29/03/2018

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

TAD Pharma GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/022/DC/18-S

Date of authorisation status change:

5/01/2022

Reference member state:

Ireland

Procedure number:

IE/V/0377/001

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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