

Simivet Retard, 150 mg/mL, suspensija za injekciju, za goveda, ovce, svinje, pse i mačke

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Simivet Retard, 150 mg/mL, suspensija za injekciju, za goveda, ovce, svinje, pse i mačke

Active substance:

Amoxicillin trihydrate

Target species:

Cattle

Sheep

Pig

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Amoxicillin trihydrate

172.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 39 day

- Milk. 108 hour

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Sheep

- Meat and offal. 29 day

VMP se ne smije primjenjivati ovcima čije se mlijeko koristi za hranu.

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Pig

- Meat and offal. 42 day

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Dog

-

Cat

Subcutaneous use:

-

Dog

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Package description:

Available only in Croatian

Available only in Croatian

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:

Dechra Regulatory B.V.

Marketing authorisation date:

13/01/2017

Manufacturing sites for batch release:

Norbrook Laboratories Limited

GENERA d.d.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/23-01/681

Date of authorisation status change:

20/10/2023

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

Documents

Summary of Product Characteristics

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