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Belacol 100 % Compactate

Authorised

- COLISTIN SULFATE

Product identification

Medicine name:

Belacol 100 % Compactate

Active substance:

COLISTIN SULFATE

Target species:

Cattle

Chicken

Pig

Route of administration:

In drinking water/milk use

Product details

Active substance and strength:

COLISTIN SULFATE

20000000.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:

Granules for use in drinking water

Withdrawal period by route of administration:**In drinking water/milk use:**

-

Cattle

- Meat and offal. 1 day

-

Chicken

- Meat and offal. 1 day

-

Pig

- Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

1 kg in fold-up carton with inner layer (paper/PE/Alu/PE)

250 g in fold-up carton with inner layer (paper/PE/Alu/PE)

2.5 kg in kard-o-seal-bag (PE/paper/PE/Alu/PE)

500 g in fold-up carton with inner layer (paper/PE/Alu/PE)

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:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

7/11/2021

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

National Organization For Medicines

Authorisation number:

105060/08-11-2021/K-0215701

Date of authorisation status change:

7/11/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0199/001

Concerned member states:

Croatia Estonia Greece Hungary Latvia Lithuania Slovakia

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

Documents

Summary of Product Characteristics

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Package Leaflet

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