Cylabel 1000 mg/g powder for use in drinking water/milk for cattle and pigs

Not authorised

- Sodium salicylate

Product identification

Medicine name:
Cylabel 1000 mg/g powder for use in drinking water/milk for cattle and pigs
Cylabel, 1000 mg/g milteliai naudoti su geriamuoju vandeniu ar pienu galvijams ir kiaulėms

Active substance:
Sodium salicylate

Target species:
Cattle (pre-ruminant)
Pig

Route of administration:
In drinking water use
In drinking water/milk use
In drinking water/milk use

Product details

Active substance and strength:
Sodium salicylate

1000.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:

In drinking water use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

In drinking water/milk use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

In drinking water/milk use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:
Lithuania

Package description:
(ID2): 1 unspecified outer container with 1 Sachet (PolyEthylene, Paper, PolyEthylene, Aluminium, PolyEthylene) with 5 kilogram(s) (5 kilogram(s))

(ID1): 1 unspecified outer container with 1 Box (Paper, PolyEthylene, Aluminium, PolyEthylene) with 1 kilogram(s) (1 kilogram(s))

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:
22/09/2017

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
SFVS

Authorisation number:
LT/2/17/2415/001-002

Date of authorisation status change:
26/01/2022

Reference member state:
Germany

Procedure number:
DE/V/0169/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents

RV2415.pdf
