Dophacyl SB 1000 mg/g, powder for use in drinking water/milk for cattle and pigs

Authorised

- Sodium salicylate

Product identification

Medicine name:
Dophacyl SB 1000 mg/g, powder for use in drinking water/milk for cattle and pigs

Active substance:
Sodium salicylate

Target species:
Cattle (calf)
Pig

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Sodium salicylate
1000.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

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

- **Cattle (calf)**
  - Meat and offal. no withdrawal period: zero days
- **Pig**
  - Meat and offal. no withdrawal period: zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BA04

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Bucket: white polypropylene square container provided with a polypropylene lid. The bucket contains 5 kg of product.
Bucket: white polypropylene square container provided with a polypropylene lid. The bucket contains 2.5 kg of product.
Bucket: white polypropylene square container provided with a polypropylene lid. The bucket contains 1 kg of product.
Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene lid. The securitainer contains 1 kg of product
- Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene lid. The securitainer contains 500 g of product

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
28/08/2023

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7009250.00.00

Date of authorisation status change:
28/08/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0392/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Spain Sweden

Generic of:
600000059248

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

Documents

Combined File of all Documents

English (PDF)
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