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Solacyl 1000 mg/g powder for use in drinking water for turkeys

Authorised

- Sodium salicylate

Product identification

Medicine name:
Solacyl 1000 mg/g powder for use in drinking water for turkeys

Active substance:
Sodium salicylate

Target species:
Turkey

Route of administration:
In drinking water 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:

- **Turkey**
  - Egg. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

  - Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BA04

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

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID6) 5 kilogram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 5 kilogram(s)
(ID5) 2.5 kilogram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 2.5 kilogram(s)
(ID4) 1 kilogram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 1 kilogram(s)
(ID3) 500 gram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 500 gram(s)
(ID2) 250 gram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 250 gram(s)
(ID1) 100 gram(s): unspecified outer container with 1 Bag (polyethylene terephthalate; aluminium; polyamide; polyethylene) with 100 gram(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:
Eurovet Animal Health B.V.

Marketing authorisation date:
31/08/2017

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402381.00.00

Date of authorisation status change:
8/11/2022

Reference member state:
Germany

Procedure number:
DE/V/0170/001

Concerned member states:
Czechia France Hungary Italy Netherlands Poland Slovakia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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