DOPHACYL AVI 1000 MG/G POWDER FOR USE IN DRINKING WATER FOR CHICKENS

Authorised

- Sodium salicylate

Product identification

Medicine name:
DOPHACYL AVI 1000 MG/G POWDER FOR USE IN DRINKING WATER FOR CHICKENS

Active substance:
Sodium salicylate

Target species:
Chicken

Route of administration:
Oral 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

Withdrawal period by route of administration:

Oral use:

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 2 weeks of the start of the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02BA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Available in:

Lithuania

Package description:

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

Sachet: white, heat sealed, 4-layer sachet with a PE inner layer which contains 100 g of product.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application - change in active substance(s) (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
11/11/2024

Manufacturing sites for batch release:
Dopharma B.V.
Dopharma France

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/24/2840/001-006

Date of authorisation status change:
10/10/2024

Reference member state:
France

Procedure number:
FR/V/0486/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia

Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Spain

---

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: 27/04/2026
Download

eu-puar-frv0486001-mr-rpe1021-en.pdf