

Phenocillin 800 mg/g powder for use in drinking water for chickens

Authorised

- Phenoxymethylpenicillin potassium

Product identification

Medicine name:

Phenocillin 800 mg/g powder for use in drinking water for chickens

Active substance:

Phenoxymethylpenicillin potassium

Target species:

Chicken

Route of administration:

Oral use

Product details

Active substance and strength:

Phenoxymethylpenicillin potassium
887.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:**Oral use:**

-

Chicken

- Eggs. 0 day

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

Bags consisting of the following materials: on the outside a polyethylene terephthalate layer, inside layers of aluminium and polyamide and an inner layer of polyethylene.

Pack sizes are 100 g.

Bags consisting of the following materials: on the outside a polyethylene terephthalate layer, inside layers of aluminium and polyamide and an inner layer of polyethylene.

Pack sizes are 10x 100 g.

Bags consisting of the following materials: on the outside a polyethylene terephthalate layer, inside layers of aluminium and polyamide and an inner layer of polyethylene.

Pack sizes are 250 g.

Bags consisting of the following materials: on the outside a polyethylene terephthalate layer, inside layers of aluminium and polyamide and an inner layer of polyethylene.

Pack sizes are 500 g.

Bags consisting of the following materials: on the outside a polyethylene terephthalate layer, inside layers of aluminium and polyamide and an inner layer of polyethylene

Pack sizes are 1000 g.

Bags consisting of the following materials: on the outside a paper layer, inside layers of polyethylene and aluminium and an inner layer of polyethylene. Pack sizes are

2500 g.

Bags consisting of the following materials: on the outside a paper layer, inside layers of polyethylene and aluminium and an inner layer of polyethylene. Pack sizes are 1000 g.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Eurovet Animal Health B.V.

Marketing authorisation date:

28/12/2015

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402170.00.00

Date of authorisation status change:

27/04/2021

Reference member state:

Ireland

Procedure number:

IE/V/0345/001

Concerned member states:

Austria Belgium Croatia Czechia Denmark France Germany Greece

Hungary Italy Lithuania Netherlands Poland Portugal Slovakia Slovenia
Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

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Package Leaflet

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