Phenoxypen WSP

Authorised

- Phenoxymethylpenicillin potassium

Product identification

Medicine name:
Phenoxypen WSP

Active substance:
Phenoxymethylpenicillin potassium

Target species:
Chicken

Route of administration:
In drinking water use
Oral use

Product details

Active substance and strength:
Phenoxymethylpenicillin potassium
325.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:

In drinking water use:

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. 0 day

Oral use:

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE02

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

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Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Securitainer: white PP cylin drical container, with white HDPE/LDPE closure, with thumb tab for opening, containing 250 g (650 ml) of product.
White polypropylene bucket provided with a polypropylene lid containing 1 kg of product.
Composite can: three layered rectangular container, which co nsists of a cardboard base with an inner lining of aluminium paper and with label on the outside containing 1 kg of product.
Securitainer: white PP cylin drical container, with white HDPE/LDPE closure, with thumb tab for opening, containing 1000 g (1875 ml) of product.

White polypropylene bucket provided with a polypropylene lid containing 2.5 kg of product.
White polypropylene bucket provided with a polypropylene lid containing 5 kg of product.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
15/03/2006

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10333

Date of authorisation status change:
25/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0121/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark France Germany Greece

Hungary Ireland Italy Lithuania Poland Portugal Romania 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

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