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Phenoxypen WSP, 325 mg/g powder for use in drinking water for pigs and chickens

Authorised

- Phenoxymethylpenicillin potassium

Product identification

Medicine name:
Phenoxypen WSP, 325 mg/g powder for use in drinking water for pigs and chickens

Active substance:
Phenoxymethylpenicillin potassium

Target species:
Chicken
Pig

Route of administration:
In drinking water use

Product details

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

Pharmaceutical form:
Powder for use in drinking water

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

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. 0 day
- **Pig**
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE02

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
White polypropylene bucket provided with a polypropylene lid containing 2.5 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 250 g (650 ml) 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 1 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:
9/01/2007

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
39520

Date of authorisation status change:
9/01/2007

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)

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Documents

Summary of Product Characteristics

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Combined File of all Documents