

Lincocin Soluble Powder, 400 mg/g Powder for Use in Drinking Water for Pigs and Chickens

Authorised

- Lincomycin hydrochloride

Product identification

Medicine name:

Lincocin Soluble Powder, 400 mg/g Powder for Use in Drinking Water for Pigs and Chickens

Active substance:

Lincomycin hydrochloride

Target species:

Chicken

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Lincomycin hydrochloride

453.63 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

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

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Chicken

- Meat and offal. 5 day

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Pig

- Meat and offal. 24 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

White high density polyethylene (HDPE) bottle containing 1.5 kg powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid. Pack sizes: Bottle of 1.5 kg

White high density polyethylene (HDPE) bottle containing 150 g powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid with an aluminium cap. Pack sizes: Bottle of 150 g

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:

21/01/1993

Manufacturing sites for batch release:

Phibro Medolla Manufacturing S.r.l.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 61652/3004

Date of authorisation status change:

1/12/2024

Reference member state:

Ireland

Procedure number:

IE/V/0410/001

Concerned member states:

Belgium France Germany Luxembourg Poland

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics