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# Lincocin Soluble Powder, 400 mg/g powder for use in drinking water

Authorised

Lincomycin hydrochloride

# Product identification

### **Medicine name:**

Lincocin Soluble Powder, 400 mg/g powder for use in drinking water Lincocin 40%, 400 mg/g poudre pour administration dans l'eau de boisson

### **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:

Luxembourg

### **Available in:**

Luxembourg

# Package description:

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

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

# 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:

Zoetis Belgium

# Marketing authorisation date:

18/06/1990

# Manufacturing sites for batch release:

Zoetis Belgium

# **Responsible authority:**

Ministry Of Health And Social Security

### **Authorisation number:**

V 087/90/10/0314

# Date of authorisation status change:

12/09/2014

### 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

English (PDF)

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