SPECTOLIKEL 50/100 mg/ml solution for injection for calves, sheep and pigs

Authorised

- Spectinomycin
- Lincomycin

Product identification

Medicine name:
SPECTOLIKEL 50/100 mg/ml solution for injection for calves, sheep and pigs
Spectoliphen 50 mg/ml Oplossing voor injectie voor kalveren, schapen en varkens
Spectoliphen 50 mg/ml Solution injectable pour veaux, ovins et porcins
Spectoliphen 50 mg/ml Injektionslösung für Kälber, Schafe und Schweine

Active substance:
Spectinomycin
Lincomycin

Target species:
Sheep
Pig
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Spectinomycin
100.00 milligram(s) / 1.00 millilitre(s)

Lincomycin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Sheep**
  - Meat and offal. 15 day

Milk: not authorised for use in animals producing milk for human consumption.

- **Pig**
  - Meat and offal. 14 day

- **Cattle**
  - Meat and offal. 23 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF52

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

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Authorisation status:
Valid

Authorised in:

Belgium

Available in:
Belgium

Package description:
250 ml translucent polypropylene vials, with bromobutyl stopper and aluminium cap with FLIP-OFF seal.
100 ml translucent polypropylene vials, with bromobutyl stopper and aluminium cap with FLIP-OFF seal.
6x250 ml translucent polypropylene vials, with bromobutyl stopper and aluminium cap with FLIP-OFF seal.
10X100 ml translucent polypropylene vials, with bromobutyl stopper and aluminium cap with FLIP-OFF seal.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
19/11/2013

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V445137

Date of authorisation status change:

19/11/2013

Reference member state:
Portugal

Procedure number:
PT/V/0111/001

Concerned member states:
Belgium

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 6/01/2026
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Package Leaflet

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Labelling

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