

LINCOMIX- lincomycin injection
Zoetis Inc.

Lincomix®

Lincomix®

brand of lincomycin injection

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

For Intramuscular Use in Swine Only

LINCOMIX Injectable contains lincomycin hydrochloride, an antibiotic produced by *Streptomyces lincolnensis* var. *lincolnensis*, which is chemically distinct from all other clinically available antibiotics and is isolated as a white crystalline solid.

INDICATIONS FOR SWINE

LINCOMIX Injectable is indicated for the treatment of infectious forms of arthritis caused by organisms sensitive to its activity. This includes most of the organisms responsible for the various infectious arthritides in swine, such as the staphylococci, streptococci, Erysipelothrix and Mycoplasma spp.

It is also indicated for the treatment of mycoplasma pneumonia.

CONTRAINDICATIONS

As with all drugs, the use of LINCOMIX Injectable is contraindicated in animals previously found to be hypersensitive to the drug.

WARNING

Swine intended for human consumption should not be slaughtered within 48 hours of latest treatment. **Not for human use.**

CAUTION

If no improvement is noted within 48 hours, consult a veterinarian.

ADVERSE REACTIONS

The intramuscular administration to swine may cause a transient diarrhea or loose stools. Although this effect has rarely been reported, one must be alert to the possibility that it may occur.

Should this occur, it is important that the necessary steps be taken to prevent the effects of dehydration.

DOSAGE AND ADMINISTRATION

For arthritis or mycoplasma pneumonia—5 mg per pound of body weight intramuscularly once daily for three to seven days as needed. When using LINCOMIX Injectable containing 100 mg/mL, 1 mL/20 lb body weight will provide 5 mg/lb. When using LINCOMIX Injectable containing 300 mg/mL, 1 mL/60 lb body weight will provide 5 mg/lb.

For optimal results, initiate treatment as soon as possible.

As with any multi-dose vial, practice aseptic techniques in withdrawing each dose. Adequately clean and disinfect the vial closure prior to entry with a sterile needle and syringe. No vial closure should be entered more than 20 times.

HOW SUPPLIED

LINCOMIX Injectable is available in two concentrations: 300 mg/mL and 100 mg/mL.

300 mg/mL: *For use in swine weighing 300 pounds or more.* Each mL contains lincomycin hydrochloride equivalent to lincomycin, 300 mg; also Benzyl Alcohol, 9.45 mg added as preservative. Supplied in 100 mL vials.

100 mg/mL: Each mL contains lincomycin hydrochloride equivalent to lincomycin, 100 mg; also Benzyl Alcohol, 9.45 mg added as preservative. Supplied in 100 mL vials.

Store at controlled room temperature 20°-25°C (68°- 77°F), with excursions between 15°- 40°C (59°- 104°F).

Use contents within 28 days of first vial broach.

Approved by FDA under NADA # 034-025

zoetis

Distributed by:
Zoetis Inc.
Kalamazoo, MI 49007

Revised: June 2022

40039342

PRINCIPAL DISPLAY PANEL - 100 mg Vial Label

For Intramuscular Use In Swine.
CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.
For Use in Animals Only.

Dosage: Usual daily dose for arthritis or mycoplasma pneumonia—5 mg per pound of body weight (1 mL per each 20 pounds of body weight) intramuscularly for three to seven days.

See package insert for complete product information.

Warnings: Not for human use.
 Keep out of the reach of children.
 Swine intended for human consumption should not be slaughtered within 48 hours of latest treatment.

Store at controlled room temperature 20°- 25°C (68°- 77°F), with excursions between 15°- 40°C (59°- 104°F).
 Use contents within 28 days of first vial broach.



Distributed by:
 Zoetis Inc., Kalamazoo, MI 49007

Lincomix® 100

Injectable

(lincomycin injection)

Swine Antibiotic



Indicated for the treatment of arthritis caused by susceptible organisms and for mycoplasma pneumonia.

Contains per mL:
 lincomycin hydrochloride equivalent to lincomycin, 100 mg; also benzyl alcohol, 9.45 mg added as preservative.

Approved by FDA under NADA # 034-025

100 mL (3.3 Fl Oz)



PRINCIPAL DISPLAY PANEL - 300 mg vial label

For Intramuscular Use In Swine Over 300 lbs.
CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.
For Use in Animals Only.

Dosage: Usual daily dose for arthritis or mycoplasma pneumonia—5 mg per pound of body weight (1 mL per each 60 pounds of body weight) intramuscularly for three to seven days.

See package insert for complete product information.

Warnings: Not for human use.
 Keep out of the reach of children.
 Swine intended for human consumption should not be slaughtered within 48 hours of latest treatment.

Store at controlled room temperature 20°- 25°C (68°- 77°F), with excursions between 15°- 40°C (59°- 104°F).
 Use contents within 28 days of first vial broach.



Distributed by:
 Zoetis Inc., Kalamazoo, MI 49007

Lincomix® 300

Injectable

(lincomycin injection)

Swine Antibiotic



Indicated for the treatment of arthritis caused by susceptible organisms and for mycoplasma pneumonia.

Contains per mL:
 lincomycin hydrochloride equivalent to lincomycin, 300 mg; also benzyl alcohol, 9.45 mg added as preservative.

Approved by FDA under NADA # 034-025

100 mL (3.3 Fl Oz)



LINCOMIX

lincomycin injection

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:54771-0617
Route of Administration	INTRAMUSCULAR		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LINCOMYCIN HYDROCHLORIDE (UNII: M6T05Z2B68) (LINCOMYCIN - UNII:BOD072YW0F)	LINCOMYCIN	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	9.45 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54771-0617-1	100 mL in 1 VIAL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA034025	06/06/1967	

LINCOMIX

lincomycin injection

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:54771-3256
Route of Administration	INTRAMUSCULAR		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LINCOMYCIN HYDROCHLORIDE (UNII: M6T05Z2B68) (LINCOMYCIN - UNII:BOD072YW0F)	LINCOMYCIN	300 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	9.45 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54771-3256-1	100 mL in 1 VIAL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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NADA

NADA034025

06/06/1967

Labeler - Zoetis Inc. (828851555)

Revised: 6/2023

Zoetis Inc.