## **SPECTINOMYCIN** (Veterinary—Systemic)

Some commonly used brand names for veterinary-labeled products are: Adspec Sterile Solution; Spectam; Spectam Injectable; Spectam Oral Solution; Spectam Scour-Halt; Spectam Powder; Spectam Water Soluble; Spectam Water Soluble Concentrate; and Spectoguard Scour-Chek.

Note: For a listing of dosage forms and brand names by country availability, see the Dosage Forms section(s).

Category: Antimicrobial (systemic).

### **Indications:**

Note: The text between ELUS and EL describes uses that are not included in U.S. product labeling. Text between ELCAN and EL describes uses that are not included in Canadian product labeling The ELUS or ELCAN designation can signify a lack of product availability in the country indicated. See the Dosage Forms

section of this monograph to confirm availability.

## **General considerations**

Spectinomycin is an antibiotic that is active against a variety of aerobic gram-negative and gram-positive organisms (R-3) as well as Mycoplasma species. (R-7) Spectinomycin is used clinically, primarily for its activity against gram-negative organisms; some gram-positive organisms may also be susceptible to this agent. It has in vitro and in vivo activity against Mannheimia (Pasteurella) haemolytica, Pasteurella multocida, and Haemophilus somnus. R-<sup>22]</sup> Anaerobic organisms are generally resistant. <sup>[R-7]</sup> Spectinomycin is usually bacteriostatic at therapeutic doses. <sup>[R-5]</sup> As an aminocyclitol antibiotic, spectinomycin is structurally and functionally similar to the aminoglycoside antibiotics, which are also aminocyclitols. Spectinomycin lacks the toxic effects of the aminoglycoside antibiotics; however, its use is limited by the ready development of bacterial resistance. {R-5}

## Accepted

Air sacculitis (treatment)<sup>EL</sup>—Turkey poults, 1- to 3-day-old: Spectinomycin hydrochloride injection is indicated to aid in the control of air sacculitis associated with Mycoplasma meleagridis sensitive to spectinomycin. (R-17)

Chronic respiratory disease (CRD) (prophylaxis)—Chickens, broiler: Spectinomycin powder for oral solution is indicated to aid in the prevention of mortality due to CRD associated with susceptible Mycoplasma gallisepticum. [R-2; 18]

Chronic respiratory disease (CRD) (treatment)—

Chickens, broiler: Spectinomycin powder for oral solution is indicated to aid in the control of mortality due to CRD associated with susceptible *Mycoplasma gallisepticum*. <sup>{R-2; 18}</sup> EL<sup>CAN</sup> *Turkey poults*, 1- to 3-day-old<sup>EL</sup>. Spectinomycin

hydrochloride injection is indicated to aid in the control of CRD associated with Escherichia coli. (R-17)

 $^{ELCAN}$ Colibacillosis (treatment) $^{EL}$ —*Chicks*, newly hatched: Spectinomycin hydrochloride injection is indicated in the control of mortality and to lessen severity of infections caused by E. coli. (R-17)

Enteritis, bacterial (treatment)—Pigs, less than four weeks of age: Spectinomycin oral solution is indicated in the treatment of bacterial enteritis (white scours) associated with *E. coli* in pigs younger than 4 weeks of age. <sup>[R.3; 4]</sup>

ELCAN Paratyphoid (treatment) EL — *Chicks*, newly hatched:

Spectinomycin hydrochloride injection is indicated in the control of mortality and to lessen severity of infections caused by Salmonella typhimurium. {R-17}

Pneumonia, bacterial (treatment)—Cattle: Spectinomycin sulfate injection is indicated in the treatment of pneumonia (bovine respiratory disease) associated with M. haemolytica, P. multocida,

and H. sommus in cattle. (R-21; 22) ELCAN Salmonella infantis infection (treatment) EL—Chicks, newly hatched: Spectinomycin hydrochloride injection is indicated in the control of mortality and to lessen severity of infections caused by S. infantis, (R-17) however, S. infantis is not considered to be a major pathogen in the poultry industry.

Synovitis (prophylaxis)—Chickens, broiler: Spectinomycin powder for oral solution is indicated to aid in the prevention of mortality associated with infectious synovitis due to susceptible *Mycoplasma synoviae*. {R-2; 18}

Synovitis (treatment)-

ELCAN Chickens, broiler EL: Spectinomycin powder for oral solution is indicated to aid in the control of mortality associated with infectious synovitis due to susceptible *M. synoviae.* <sup>{R-18}</sup>

ELCAN Chicks, newly hatched EL: Spectinomycin hydrochloride injection is indicated in the control of mortality and to lessen severity of infections caused by susceptible M. synoviae. {R-17}

ELUS Fowl cholera (treatment)EL—Turkeys: Spectinomycin hydrochloride injection is indicated to reduce mortality due to fowl cholera caused by sensitive strains of Pasteurella  $multocida.^{\{R-1\}}$ 

## Potentially effective

Colibacillosis (treatment)<sup>EL</sup>—ELUS Ducklings<sup>EL</sup>: For use in animals not to be used in the production of human food—There are insufficient data to establish the safety and efficacy of spectinomycin in the treatment of colibacillosis in ducklings; however, in one study, subcutaneous administration of spectinomycin reduced the mortality and improved weight gain in 1-day-old ducklings experimentally infected with *E. coli.* (R-10)

Infections, bacterial (treatment), including Respiratory tract infections (treatment)—ELUS,CAN PigsEL: There are insufficient data to establish the safety and efficacy of spectinomycin injection in the treatment of respiratory infections and systemic infections due to susceptible organisms in pigs; however, the parenteral administration of spectinomycin to pigs has been used in clinical practice to treat these infections. (R

## **Regulatory Considerations**

Spectinomycin is not labeled for use in birds producing eggs for human consumption. (R-18)

Withdrawal times have been established for the use of spectinomycin in newly hatched chicks, [R-17] broiler chickens, (R-18) 1- to 3-day-old turkey poults, (R-17) and piglets (R-4) (see the *Dosage Forms* section).

Canada-

Spectinomycin is not labeled for use in birds producing eggs for human consumption. (R-1; 2)

Spectinomycin injection is not labeled for use in turkeys weighing <0.5 mg. (R-1)

Withdrawal times have been established for the use of spectinomycin in broiler chickens, {R-2} piglets, {R-3} and turkeys. (see the *Dosage Forms* section).

## Chemistry

Source: Spectinomycin is a product of Streptomyces spectabilis. (R-5;

Chemical group: Aminocyclitol. [R-5] Chemical name:

Spectinomycin hydrochloride—4H-Pyrano[2,3b][1,4]benzodioxin-4-one, decahydro-4a,7,9-trihydroxy-2methyl-6,8-bis(methylamino)-, dihydrochloride, pentahydrate. [R-6]

Spectinomycin sulfate tetrahydrate—Decahydro-4a,7,9trihydroxy-2-methyl-6,8-bis(methylamino)-4H-pyrano[2,3b][1,4]benzodioxin-4-one sulfate, tetrahydrate. {R-22}

Molecular formula: Spectinomycin hydrochloride—

 $C_{14}H_{24}N_2O_7 \cdot 2HCl \cdot 5H_2O.^{\{R-6\}}$ 

**Molecular weight:** Spectinomycin hydrochloride—495.35.<sup>[R-6]</sup> **Description:** Spectinomycin Hydrochloride USP—White to palebuff crystalline powder.<sup>[R-16]</sup>

**pKa:** 6.95 and 8.70. {R-19}

**Solubility:** Spectinomycin Hydrochloride USP—Freely soluble in water; practically insoluble in alcohol, in chloroform, and in ether. (R-16)

## Pharmacology/Pharmacokinetics

Note: Unless otherwise noted, pharmacokinetic data in this section are based on a single intravenous injection of spectinomycin.

The pharmacokinetics and detection of spectinomycin do not appear to be influenced by administration in combination with lincomycin; (R-7) some of the pharmacokinetic data in this section are derived from studies in which lincomycin and spectinomycin were administered concomitantly. (R-7)

**Mechanism of action/Effect:** Spectinomycin binds to the 30S ribosomal subunit of the microorganism and inhibits protein synthesis by preventing elongation of the polypeptide chain at the translocation step. [R-5]

Absorption: Spectinomycin is only slightly absorbed from the gastrointestinal tract; (R-7) however, it is rapidly absorbed following intramuscular administration. (R-7) In cattle, spectinomycin is completely bioavailable following intramuscular administration. (R-7) Repeated administration in cattle does not appear to result in tissue concentrations higher than those achieved with a single dose. (R-7)

Distribution: Twelve hours following intramuscular administration and 24 hours following oral administration, concentrations of spectinomycin are found in the following swine tissues in decreasing concentrations: kidney, liver, lung, muscle, and fat. (R-7) An identical profile is seen in cattle 24 and 72 hours following intramuscular administration of spectinomycin. (R-7) Tissue/serum ratios of spectinomycin usually do not exceed 0.25 to 0.5 and are much lower in brain, aqueous humor, and bone.

Volume of distribution (Vol<sub>D</sub>):

Cows—0.295 Liter per kg (L/kg). {R-13} Ewes—0.307 L/kg. {R-13}

Protein binding: Cows—Low (approximately 10%). [R-13]

**Biotransformation:** Spectinomycin does not appear to undergo any significant metabolism. In swine, it is excreted unchanged in the urine following intramuscular administration. <sup>[R-7]</sup>

Half-life: Elimination—

Cows: 1.01 to 1.2 hours. (R-7; 13) Ewes: 1.01 hours. (R-13) Pigs: 0.98 hour. (R-7)

### **Peak serum concentration:**

Calves, preruminating—20 micrograms per mL (mcg/mL) between 0.33 and 0.67 hours following an intramuscular dose of 10 mg per kg of body weight (mg/kg). [R-7]

Cows—Approximately 55 mcg/mL at 1 hour following an intramuscular dose of 20 mg/kg. (R-13)

Dogs-

Intramuscular: 78 mcg/mL 40 minutes following an intramuscular dose of 40 mg/kg.

Oral:<sup>{**R**-7</sup>

22 mcg/mL approximately 4 hours following a dose of 100 mg/kg.

80 mcg/mL approximately 4 hours following a dose of

500 mg/kg.

Ewes—Approximately 53 mcg/mL at 1 hour following an intramuscular dose of 20 mg/kg.<sup>[R-13]</sup>

#### **Elimination:**

Following intramuscular administration—Spectinomycin is rapidly absorbed, then quickly eliminated from plasma and tissues through renal excretion. (R-7) Because of this rapid excretion, drug accumulation is not observed following repeated administration. (R-7) Renal impairment may cause accumulation of the active drug.

Following oral administration—Because spectinomycin is poorly absorbed from the gastrointestinal tract, it is excreted mostly in the feces. (R-7)

## **Precautions to Consider**

## Lactation

Cows: In one experimental study, the milk-to-serum ratio of spectinomycin concentrations ranged from 0.44 to 1.12 in mastitic cows receiving one intramuscular dose of 20 mg per kg of body weight (mg/kg), followed by three intramuscular doses of 10 mg/kg at hourly intervals. [R-13] Spectinomycin levels in milk from dairy cows receiving an intramuscular dose of 20 mg/kg two times a day for 3 consecutive days were below 0.2 mcg/mL at the fifth milking after the last injection. [R-7] No residues of spectinomycin were detectable at the seventh milking. [R-7]

## **Side/Adverse Effects**

The following side/adverse effects have been selected on the basis of their potential clinical significance (possible signs and, for humans, symptoms in parentheses where appropriate)—not necessarily inclusive:

## Those indicating need for medical attention

Incidence unknown

All species

Anaphylactic reactions; (R-22) neuromuscular blockade (R-5)

## Those indicating need for medical attention only if they continue or are bothersome

Incidence unknown

Cattle

Discoloration of tissue at the injection site;  $^{\{R-22\}}$  swelling at the injection site, mild $^{\{R-22\}}$ 

## Human side/adverse effects<sup>{R-15}</sup>

In addition to the above side/adverse effects reported in animals, the following side/adverse effects have been reported in humans, and are included in the human monograph *Spectinomycin (Systemic)* in *USP DI Volume I;* these side/adverse effects are intended for informational purposes only and may or may not be applicable to the use of spectinomycin in the treatment of animals: Incidence rare

Dizziness; gastrointestinal disturbance; hypersensitivity; pain at site of injection

## **Overdose**

For more information in cases of overdose or unintentional ingestion, contact the American Society for the Prevention of Cruelty to Animals (ASPCA) National Animal Poison Control Center (888-426-4435 or 900-443-0000; a fee may be required for consultation) and/or the drug manufacturer.

Cattle: When cattle were administered 150 mg per kg a day (10 times the labeled dose) for 5 days, the effects seen at the end of the treatment period included increased relative kidney weights. {R-22} Urinalysis was performed only on steers. Urinary pH was decreased and squamous and transitional cells were found in the urine. {R-22}

### Clinical effects of overdose

Note: The following effects have been selected on the basis of their potential clinical significance (possible signs in parentheses where appropriate)—not necessarily inclusive (» = major clinical significance):

Acute effects-

 $Turkey\ poults^{\{R-1\}}$ 

#### Ataxia: coma

Note: Clinical signs of *ataxia* and *coma* following a single, subcutaneous dose of 90 mg per poult were transient, resolving after 4 hours; (R-1) a single, subcutaneous injection of up to 50 mg per poult caused no detectable ill effects. (R-1)

## **Veterinary Dosing Information**

## Safety considerations

Some individuals who handle spectinomycin develop serious reactions involving skin, nails, and eyes. (R-1; 9) Individuals who have experienced a rash or other evidence of allergic reaction should avoid further contact with spectinomycin. (R-2)

## **Oral Dosage Forms**

Note: The dosing and strengths of the dosage forms available are expressed in terms of spectinomycin base (not the hydrochloride salt).

The text between ELUS and EL describes uses not included in U.S. product labeling. Text between ELCAN and EL describes uses that are not included in Canadian product labeling.

The  $^{\mathrm{ELUS}}$  or  $^{\mathrm{ELCAN}}$  designation can signify a lack of product availability in the country indicated. See also the Strength(s) usually available section for each dosage form.

# SPECTINOMYCIN HYDROCHLORIDE ORAL SOLUTION

**Usual dose:** Enteritis, bacterial—*Pigs*, younger than 4 weeks of age or less than 6.8 kg of body weight:

For pigs weighing < 4.5 kg—Oral, 50 mg (base) as a total dose per animal two times a day for three to five days. (R-3; 4)

For pigs weighing 4.5 kg to 6.8 kg—Oral, 100 mg (base) as a total dose per animal two times a day for three to five days. [R-3; 4] Withdrawal times—US and Canada: Meat—21 days. [R-3; 4]

Note: If improvement is not seen within forty-eight hours of initiating treatment, the diagnosis or choice of therapy should be reconsidered. (R-3; 4)

## Strength(s) usually available:

U.S.-

For veterinary-labeled product(s):

50 mg (base) per mL (OTC) [Spectam Scour-Halt; Spectoguard Scour-Chek].

Canada-

For veterinary-labeled product(s):

50 mg (base) per mL (OTC) [Spectam Oral Solution; Spectam Scour-Halt].

Packaging and storage: Store at controlled room temperature, 20 to 25 °C (68 to 77 °F). <sup>[R-4]</sup> Do not freeze. <sup>[R-3]</sup>

**Auxiliary labeling:** When not in use, the plastic doser should be removed and the original cap replaced on bottle. [R-3; 4] The plastic doser should be rinsed with water after each use.

USP requirements: Not in USP. [R-16]

# SPECTINOMYCIN HYDROCHLORIDE POWDER FOR ORAL SOLUTION

## Usual dose:

Chronic respiratory disease (prophylaxis and control)—Chickens,

broiler: Oral, administered as the sole source of drinking water at a concentration of 0.5 mg (base) per mL (2 grams [base] per gallon) of water for the first three days of life and for one day following each vaccination. (R-2; 18)

for one day following each vaccination. (R-2; 18)

Synovitis (prophylaxis and ELCAN treatment EL)—Chickens, broiler:

Oral, administered as the sole source of drinking water at a concentration of 0.26 mg (base) per mL (1 gram [base] per gallon) of water for the first three to five days of life. (R-18)

Note: Canadian labeling lists a dose of 0.5 mg (base) per mL (2 grams [base] per gallon) of water for this indication. (R-2)

Withdrawal times—US and Canada: Meat—5 days. <sup>{R-2; 18; 19}</sup>
Products are not labeled for use in poultry laying eggs for human consumption. <sup>{R-18}</sup>

## Strength(s) usually available:

U.S.-

Veterinary-labeled product(s):

500 mg (base) per gram of water-soluble powder (OTC) [Spectam Water Soluble].

Canada-

Veterinary-labeled product(s):

500 mg (base) per gram of water-soluble powder (OTC) [Spectam Powder; Spectam Water Soluble Concentrate].

**Packaging and storage:** Store below 40 °C (104 °F), preferably between 15 and 30 °C (59 and 86 °F), unless otherwise specified by manufacturer.

Preparation of dosage form: Water-soluble powder should be mixed with drinking water according to the manufacturer's directions

USP requirements: Not in USP. {R-16}

## **Parenteral Dosage Forms**

Note: The dosing and strengths of the dosage forms available are expressed in terms of spectinomycin base (not the hydrochloride or sulfate salt)

The text between <sup>ELUS</sup> and <sup>EL</sup> describes uses not included in U.S. product labeling. Text between <sup>ELCAN</sup> and <sup>EL</sup> describes uses that are not included in Canadian product labeling. The <sup>ELUS</sup> or <sup>ELCAN</sup> designation can signify a lack of product

The ELUS or ELCAN designation can signify a lack of product availability in the country indicated. See also the *Strength(s)* usually available section for each dosage form.

# SPECTINOMYCIN HYDROCHLORIDE INJECTION Usual dose:

ELCAN Air sacculitis (treatment) EL — Turkey poults, 1- to 3-day-old:

Subcutaneous in cervical area, 10 mg (base) as a single, total dose per poult. [R-17]

Withdrawal times—US: Meat—0 days. {R-17}

ELCAN Chronic respiratory disease (treatment) EL—Turkey poults, 1to 3-day-old: Subcutaneous in cervical area, 5 mg (base) as a single, total dose per poult. (R-17) Dilution with sterile physiologic saline is recommended to facilitate accurate dosing. (R-17)

Withdrawal times—US: Meat—0 days. {R-17}

ELCAN Colibacillosis (treatment)EL;

ELCAN Paratyphoid (treatment)EL;

ELCAN Salmonella infantis infection (treatment)<sup>EL</sup>; or

ELCAN Synovitis (treatment) EL—Chicks, newly hatched: Subcutaneous in cervical area, 2.5 to 5 mg (base) as a single, total dose per chick. (R-17) Dilution with sterile physiologic saline is recommended so that the total volume administered is 0.2 mL. (R-17)

Withdrawal times—US: Meat—0 days. {R-17}

FLUS Fowl cholera (treatment) FL—Turkeys: Subcutaneous in dorsal

cervical area, 11 to 22 mg (base) per kg of body weight as a single injection. The entire flock should be treated as soon as symptoms of fowl cholera are seen.<sup>{R-1}</sup> Treatment must not be repeated within five days of the initial treatment.<sup>{R-1}</sup> Withdrawal times—Canada: Meat—5 days.<sup>{R-1}</sup>

Note: ELUS,CAN Ducklings EL—For use in animals not to be used in the production of human food: Although there are insufficient data to establish safety and efficacy, a single, subcutaneous, total dose of 5 mg (base) per duckling has been shown to reduce mortality and improve weight gain in one-day-old ducklings experimentally infected with E. coli. [8-10]

ELUS,CAN PigsEL—Although there are insufficient data to establish safety and efficacy, the intramuscular administration of spectinomycin to pigs, at doses ranging from 6.6 to 22 mg (base) per kg of body weight every twelve to twenty-four hours, [R-11] has been used in clinical practice to treat respiratory infections and systemic infections caused by organisms sensitive to spectinomycin. [R-5]

Extra-label withdrawal recommendation: U.S. and Canada—*Pigs*: Because injectable spectinomycin is not labeled for use in pigs, there are no established withdrawal times in the U.S. or Canada. If spectinomycin is administered intramuscularly at a dose of 20 mg per kg of body weight, evidence has been compiled by the Food Animal Residue Avoidance Databank (FARAD) and Canadian gFARAD that suggests a meat withdrawal interval of thirty days would be sufficient to avoid violative residues. <sup>{R-7; 14; 23}</sup>

## Strength(s) usually available:

HS —

Veterinary-labeled product(s):

100 mg (base) per mL (OTC) [Spectam Injectable]. [R-17]
Canada—

Veterinary-labeled product(s):

100 mg (base) per mL (OTC) [Spectam Injectable]. [R-1]

## Preparation of dosage form:

Dilution with sterile physiologic saline according to product labeling is recommended when administering total doses ≤ 5 mg and is appropriate when large flocks are being treated. (R-17) Aseptic technique must be employed and unused diluted solution should be discarded. (R-17)

Packaging and storage: Store at controlled room temperature 20 to 25 °C (68 to 77 °F), unless otherwise specified by the manufacturer. Protect from freezing. (R-17)

Auxiliary labeling: Injection site should be disinfected prior to injection and precautions should be taken to prevent contamination of the contents of the bottle. (R-1; 17)

USP requirements: Not in USP. (R-16)

## SPECTINOMYCIN SULFATE INJECTION

**Usual dose:** Pneumonia—*Cattle:* Subcutaneous, 10 to 15 mg (base) per kg of body weight every twenty-four hours for three to five days. [R-22]

Withdrawal times—US and Canada: Meat—11 days. <sup>{R-21; 22}</sup> Product labeling states that withdrawal times have not been established for lactating dairy cattle. <sup>{R-21; 22}</sup> US product labeling states that a withdrawal period has not been established for preruminating calves. <sup>{R-22}</sup> Discoloration of tissue at the injection site may last more than 11 days, making it necessary to trim the site and surrounding tissue at slaughter <sup>{R-22}</sup>; Canadian product labeling recommends not slaughtering cattle for at least 15 days to avoid excessive trimming. <sup>{R-21}</sup>

Note: It is recommended that this medication be administered

subcutaneously in the neck and that not more than 50~mL be given per site. (R-22)

## $Strength(s) \ usually \ available; \ ^{\{R-21;\ 22\}}$

U.S.—

Veterinary-labeled product(s):

100 mg (base) per mL (Rx) [Adspec Sterile Solution].

Veterinary-labeled product(s):

100 mg (base) per mL (Rx) [Adspec Sterile Solution].

Package and storage: Store at 20 to 25 °C (68 to 77 °F), unless otherwise specified by the manufacturer. (R-22) Protect from freezing.

USP requirements: Not in USP. {R-16}

Developed: 07/08/98

Interim revision: 10/15/99; 09/30/02; 04/05/03

Revision: 09/29/08

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