SULFAMED- sulfadimethoxine injection Bimeda, Inc.

SulfaMed™ (sulfadimethoxine) Injection INJECTION 40% ANTIBACTERIAL Each mL contains 400 mg sulfadimethoxine

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

Cattle: For the treatment of bovine respiratory disease complex (shipping fever complex) and bacterial pneumonia associated with *Pasteurella* spp.sensitive to sulfadimethoxine; necrotic pododermatitis (foot rot) and calf diphtheria caused by *Fusobacterium necrophorum*sensitive to sulfadimethoxine.

Restricted Drug (California) - Use Only as Directed

Not for Human Use

DESCRIPTION: SulfaMed (sulfadimethoxine) Injection 40% is a low-dosage, rapidly absorbed, long-acting sulfonamide, effective for the treatment of shipping fever complex, bacterial pneumonia, calf diphtheria, and foot rot in cattle. Sulfadimethoxine is a white, almost tasteless and odorless compound. Chemically, it is N1-(2,6-dimethoxy-4-pyrimidinyl) sulfanilamide. The structural formula is:



ACTIONS: Sulfadimethoxine has been demonstrated clinically or in the laboratory to be effective against a variety of organisms, such as streptococci, klebsiella, proteus, shigella, staphylococci, escherichia, and salmonella. ^{1,2}

The systemic sulfonamides which include sulfadimethoxine are bacteriostatic agents. Sulfonamides competitively inhibit bacterial synthesis of folic acid (pteroylglutamic acid) from para-aminobenzoic acid. Mammalian cells are capable of utilizing folic acid in the presence of sulfonamides.

The tissue distribution of sulfadimethoxine, as with all sulfonamides, is a function of plasma levels, degree of plasma protein binding, and subsequent passive distribution in the tissues of the lipid-soluble unionized form. The relative amounts are determined by both its pKa and by the pH of each tissue. Therefore, levels tend to be higher in less acid tissue and body fluids or those diseased tissues having high concentrations of leucocytes.²

Slow renal excretion results from a high degree of tubular reabsorption,³ and plasma protein binding is very high, providing a blood reservoir of the drug. Thus, sulfadimethoxine maintains higher blood levels than most other long-acting sulfonamides. Single, comparatively low doses of sulfadimethoxine give rapid and sustained therapeutic blood levels.¹

To assure successful sulfonamide therapy (1) the drug must be given early in the course of the disease, and it must produce a high sulfonamide level in the body rapidly after administration, (2) therapeutically effective sulfonamide levels must be maintained in the body throughout the treatment period, (3) treatment should continue for a short period of time after the clinical signs have disappeared, and (4) the causative organisms must be sensitive to this class of drugs.

TOXICITY AND SAFETY: Data regarding acute (LD50) and chronic toxicities of sulfadimethoxine indicate the drug is very safe. The LD50 in mice is greater than 2 g/kg of body weight when administered intraperitoneally and greater than 16 g/kg when administered orally. In dogs receiving massive single oral doses of 3.2 g/kg of body weight, diarrhea was the only adverse effect observed. Dogs given 160 mg/kg of body weight orally daily for 13 weeks showed no signs of toxicity.

In cattle sulfadimethoxine has been shown to be safe through extensive clinical use with other dosage forms. In addition, studies with intravenous administration of sulfadimethoxine injection 40% have demonstrated that hemolysis of erythrocytes does not occur by this route of administration. Sulfadimethoxine has a relatively high solubility at the pH normally occurring in the kidney, precluding the possibility of precipitation and crystalluria.

INDICATIONS: SulfaMed Injection 40% is indicated for the treatment of bovine respiratory disease complex (shipping fever complex) and bacterial pneumonia associated with *Pasteurellaspp*. sensitive to sulfadimethoxine; necrotic pododermatitis (foot rot) and calf diphtheria caused by *Fusobacterium necrophorum*sensitive to sulfadimethoxine.

LIMITATIONS: Sulfadimethoxine is not effective in viral or rickettsial infections, and as with any antibacterial agent, occasional failures in therapy may occur due to resistant microorganisms. The usual precautions in sulfonamide therapy should be observed.

WARNING: Milk taken from animals during treatment and for 60 hours (5 milkings) after the latest treatment must not be used for food. Do not administer within 5 days of slaughter.

A withdrawal period has not been established for this product in preruminating calves.

DO NOT USE IN CALVES TO BE PROCESSED FOR VEAL.

PRECAUTIONS: During treatment period, make certain that animals maintain adequate water intake. If animals show no improvement within 2 or 3 days, consult your veterinarian.

Tissue damage may result from perivascular infiltration.

DOSAGE AND ADMINISTRATION: SulfaMed Injection 40% must be administered only by the intravenous route in cattle. Cattle should receive 1 mL of SulfaMed Injection 40%

per 16 lb of body weight (55 mg/kg) as an initial dose, followed by 0.5 mL per 16 lb of body weight (27.5 mg/kg) every 24 hours thereafter. Sulfadimethoxine Boluses may be utilized for maintenance therapy in cattle. Representative weights and doses are indicated in the following table:

Animal Weight	Initial Dose 25 mg/lb (55 mg/kg)	Subsequent Daily Doses 12.5 mg/lb (27.5 mg/kg)		
250 lb (113.6 kg)	15.6 mL	7.8 mL		
500 lb (227.2 kg)	31.2 mL	15.6 mL		
750 lb (340.9 kg)	46.9 mL	23.5 mL		
1000 lb (454.5 kg)	62.5 mL	31.3 mL		

Each mL contains 400 mg sulfadimethoxine

Length of treatment depends on the clinical response. In most cases treatment for 3 - 5 days is adequate. Treatment should be continued until the animal is asymptomatic for 48 hours.

DIRECTIONS FOR INTRAVENOUS INJECTION: Equipment Needed:

 A nose lead and/or halter sufficiently strong enough to effectively restrain or hold the animal's head steady so that the intravenous injection can be made with ease.
Hypodermic needles, 16 or 18 gauge and 2 inches long. Only new, sharp and sterile hypodermic needles should be used. Dull needles should be discarded. Extra needles should always be available in case the needle being used should become clogged.
Hypodermic syringes, 40- or 50-mL sterile, disposable or reusable glass syringes should be available.

4. Alcohol (70%) or equally effective antiseptic for disinfecting the skin.

Preparation of equipment: Glass syringes and regular hypodermic needles should be thoroughly cleaned and washed. Following this, the needles and syringes should be immersed in boiling water for 30 minutes prior to each injection. Regular hypodermic needles should not be used more than 3-4 times as repeated skin puncturing and boiling of the needles causes them to become quite dull. Disposable hypodermic needles and syringes should not be used more than once.

Restraint of animal: The cow should preferably be in a stanchion for maximum restraint. If this is not possible, the animal should be restrained in a manner to prevent excessive movement. A nose lead should be applied and the animal's head turned sidewise to stretch the skin and tense the muscles of the neck region (See Figure 1).



Locating the jugular vein: Once the animal has been restrained (as above), you will notice a long depression of the skin from below the angle of the jaw to just above the shoulder. This is known as the jugular furrow or jugular groove. The jugular vein is located just under the jugular groove (See Figure 1).

Preparation of SulfaMed Injection 40% for injection: The rubber cap of the bottle should be thoroughly cleaned with 70% alcohol or other satisfactory antiseptic. The correct amount of SulfaMed Injection 40% for treatment should be calculated (see dosage directions) and that amount withdrawn into a syringe (See Figure 2). One or two syringefuls of air should be injected into the bottle first to make withdrawing the drug easier. SulfaMed Injection 40% should preferably be at room temperature when filling syringes and when injecting intravenously.



Entering the vein: The skin of the injection area should be clean and free of dirt. Cotton saturated with 70% alcohol (or suitable antiseptic) should be used to wipe the injection site. Apply pressure over the jugular vein close to the shoulder. This will reduce the flow of blood to the heart and cause the jugular vein to bulge or enlarge (See Figure 3).



When the jugular vein has been "raised", insert the hypodermic needle at a 45° angle through the skin just underneath the jugular vein. The beveled edge of the hypodermic needle should be up (See Figure 4).



After the skin has been punctured, the point of the needle should be directed toward the side of the vein and pushed into the center of the vein (See Figure 5).



When the needle is in the center of the vein, there will be free flow of blood back through the needle. Release external pressure when you are sure the needle is within the vein.

Injecting the SulfaMed Injection 40%: After the needle has been accurately inserted into the jugular vein, firmly attach the syringe containing SulfaMed Injection 40% to the inserted hypodermic needle. Caution; be sure syringe is free of air. Exert firm pressure on the plunger of the syringe to inject the SulfaMed Injection 40% while the barrel is held firmly. The injection should be made moderately slow - never rapidly. If the animal moves, causing resistance in pushing the plunger of the syringe, or if a bubble of the drug is noted under the skin, the needle is no longer within the vein. The needle should be repositioned.

When the injection is completed, quickly withdraw the syringe and needle with a quick pull and apply light pressure over the injection site with alcohol and cotton to minimize bleeding from the puncture site.

To report suspected adverse drug events, for technical assistance or to obtain a copy of the Safety Data Sheet (SDS), contact Bimeda, Inc. at 1-888-524-6332. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or online at www.fda.gov/reportanimalae.

STORAGE: Store at 20°C - 25°C (68°F - 77°F); excursions permitted between 15°C - 30°C (between 59°F - 86°F).

HOW SUPPLIED: SulfaMed Injection 40% is available in the following dosage form for cattle: SulfaMed Injection 40% - Each mL contains 400 mg sulfadimethoxine compounded with 20% propylene glycol, 1% benzyl alcohol, 0.1 mg disodium edetate, 1 mg sodium formaldehyde sulfoxylate, and pH adjusted with sodium hydroxide, 250 mL bottles.

REFERENCES

1. Data on file from Hoffman-La Roche Inc., Nutley, New Jersey.

Stowe CM: The sulfonamides. In Jones LM (ed). *Veterinary Pharmacology and Therapeutics*, Ames, Iowa, Iowa State University Press, chapter 33, 1965.
Baggot, JD: Some aspects of drug persistence in domestic animals. Res Vet Sci 11:(2) 130, 1970.



SULFAMED					
sulfadimethoxine injection					
Product Information					
Product Type	PRESCRIPTION ANIMAL DRUG	ltem Co	de (Source)	NDO	C:61133-6016
Route of Administration	INTRAVENOUS				
	N/a : a to a				
Active Ingredient/Active	моюту				
Ingredient Name Basis of Strength					Strength
SULFADIMETHOXINE (UNII: 30CPC UNII: 30CPC5LDEX)	C5LDEX) (SULFADIMETHOXINE -		SULFADIMETHOXI	NE	400 mg in 1 mL
Inactive Ingredients					
	Ingredient Name				Strength
PROPYLENE GLYCOL (UNII: 6DC90	Q167V3)				
BENZYL ALCOHOL (UNII: LKG8494WBH)					
EDETATE DISODIUM (UNII: 7FLD91C86K)					
SODIUM FORMALDEHYDE SULFO	DXYLATE (UNII: X4ZGP7K714)				
SODIUM HYDROXIDE (UNII: 55X04	QC32I)				

Packaging						
#	ltem Code	Package Description	Marketing	Start Date	Marke	eting End Date
1	NDC:61133-6016-1	250 mL in 1 BOTTLE				
Marketing Information						
	Marketing	Application Number or Mo	onograph	Marketing S	tart	Marketing End
	Category	Citation		Date		Date
٨N	ADA .	ANADA200523		05/15/2014		

Labeler - Bimeda, Inc. (060492923)

Establishment

Name	Address	ID/FEI	Business Operations
Bimeda-MTC Animal Health		256232216	manufacture

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Bimeda, Inc.