

RESTRICTED VETERINARY MEDICINE

Keep out of reach of children

FOR ANIMAL TREATMENT ONLY

Tribrissen[®] 48%

ANTIBIOTIC

Each mL contains 80mg trimethoprim and 400mg sulphadiazine in an aqueous suspension.

50mL

Read entire label before use

HANDLING PRECAUTIONS WARNING May be harmful if swallowed or absorbed through the skin. May cause mild skin and eye irritation. May cause an allergic skin reaction. May cause bone damage from repeated oral exposure at high doses. Avoid skin and eye contact. Wear gloves and wash thoroughly after handling. Do not breathe vapours.

FIRST AID If swallowed immediately call the National Poisons Centre 0800 POISON (0800 764 766) or a doctor. Do NOT induce vomiting. **If on skin** wash with plenty of soap and water. If skin irritation occurs or you feel unwell: get medical advice. **If in eyes** rinse cautiously with water for several minutes. If eye irritation persists, get medical advice. **If inhaled** remove to fresh air.

DISPOSAL Dispose of unused contents in a suitable landfill. Dispose of empty container by burying in a suitable landfill, or if appropriate recycle. Avoid contamination of any water supply with product or empty container.

EMERGENCY RESPONSE For specialist advice in an emergency call 0800 CHEMCALL (0800 243 622)

See Safety Data Sheet for further information. www.intervet.co.nz.

USES

Tribrissen 48% injection is indicated for the treatment and control of a wide range of respiratory, gastrointestinal and urogenital tract infections caused by sensitive organisms in cattle, sheep, goats, pigs and horses. Other infections such as foot infections, severe mastitis, bacterial agalactia of sows, and infections of the eye, ear and mouth may also be susceptible.

Contraindications

Do not administer i/m or s/c in horses, ponies and foals because of the potential for injection site reactions. Potentiated sulphonamides should not be administered by intravenous injection to anaesthetised or sedated horses as potentially fatal cardiac dysrhythmias can occur as a consequence.

Not for use in bobby calves.

DOSAGE

16mg/kg (i.e. 1mL/30kg bodyweight daily)

Horses:	slow i/v only	1mL/30kg
Cattle, sheep, goats and pigs:	i/m or slow i/v	1mL/30kg

In cases of severe infection the dose may be increased to 24mg/kg (ie: 1.5mL/30kg) bodyweight daily. A single injection may be sufficient in uncomplicated conditions but in severe or complicated infections the dose should be repeated daily for five days or until two days after the symptoms are no longer apparent.

ADMINISTRATION

Shake well before use.

Administer in the anterior half of the neck in food producing animals.

Horses: Administer by slow intravenous injection only.

Cattle, sheep, goats and pigs: Administer by intravenous or deep intramuscular injection.

WITHHOLDING PERIOD

It is an offence for users of this product to cause residues exceeding the relevant MRL in the New Zealand (Maximum Residues Limits of Agricultural Compounds) Food Standards.

Meat: Cattle, sheep, goats and pigs producing meat or offal for human consumption must not be sold for slaughter either during treatment or within 28 days of the last treatment. **Horses** 63 days.

Milk: Cattle Milk intended for sale for human consumption must be discarded during treatment and for not less than 48 hours (4 milkings) following the last treatment. **Sheep and goats** 35 days.

STORAGE

Store away from light at 2°C to 25°C.

ACVM Registration No. A5320.

See www.nzfsa.govt.nz/acvm/ for registration conditions.

Registered pursuant to the HSNO Act 1996, HSR002215

See www.ermanz.govt.nz for full controls.

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Registered to:

Schering-Plough Animal Health Limited

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