

For Animal Use Only

NOROTRIM 24 Injection

Reg. No G 2620 (Act 36/1947)

For use by, or under the control of a veterinarian only.

Storage:

Store in a cool dry place. Protect from light.

Composition:

Sulphadiazine 200,0 mg and Trimethoprim 40,0 mg per ml.

Indications for Use:

Norotrim 24 is indicated for the treatment of pneumonia, bacterial scours, urogenital tract infections, footrot, joint-ill and other infections caused by sulphonamide-sensitive organisms in horses, cattle, sheep, goats, pigs, dogs and cats. Its spectrum covers both Gram-positive and Gram-negative bacteria including: *Actinobacillus spp.*, *Actinomyces bovis*, *Bordetella spp.*, *Corynebacterium spp.*, *Escherichia coli*, *Fusiformis spp.*, *Haemophilus spp.*, *Klebsiella spp.*, *Listeria monocytogenes*, *Nocardia spp.*, *Pasteurella spp.*, *Proteus spp.*, *Salmonella spp.*, *Staphylococcus spp.* and *Streptococcus spp.*

Norotrim 24 is also indicated for the treatment of respiratory infections of bacterial origin such as rhinitis and bronchitis as well as other gastro-intestinal tract infections.

Warnings:

Milk should not be used for human consumption until 48 hours after the last treatment. Cattle should not be slaughtered for human consumption until 12 days after last treatment. Pigs should not be slaughtered for human consumption until 15 days after last treatment. Sheep should not be slaughtered for human consumption until 28 days after last treatment.

Do not use in animals that are sensitive to sulphonamides, have severe liver damage or blood dyscrasias. Do not use in horses exhibiting drug-induced cardiac arrhythmias, these may be associated with certain anaesthetics or sedatives.

Do not give the injections by routes other than those recommended. Do not inject intraperitoneally. Intravenous injections should be administered slowly in order to prevent anaphylactic shock. Ensure that adequate drinking water is available during the treatment period.

Keep out of reach of children, uninformed persons and animals.

Although this remedy has been extensively tested under a large variety of conditions, failure thereof may ensue as a result of a wide range of reasons. If this is suspected, seek veterinary advice and notify the registration holder.

Precautions:

This product may crystallize at low temperatures. These may be dissolved by gently warming.

Directions for Use:

Use only as directed.

Horses, cattle, sheep, goats and pigs:

1 ml per 16 kg bodymass daily by intramuscular or slow intravenous injections. In cases of severe infection this may be increased to 1 ml per 10 kg bodymass daily.

Dogs and Cats:

1 ml per 8 kg bodymass daily by subcutaneous injection only. The recommended site in dogs is under the loose skin at the back of the neck.

A single injection may be sufficient in uncomplicated cases, but in severe infections the dosage may be repeated for up to 5 days.

Registration Holder:

Biotech Laboratories (Pty) Ltd
Ground Floor, Block K West,
Central Park
400 16th Road, Midrand, Gauteng, 1685

Marketed by:

Biotech Laboratories (Pty) Ltd
Tel: (011) 848 3050
Fax: (011) 848 3065
Email: info@biotechlabs.co.za
www.biotechlabs.co.za

Slegs Vir Dieregebruik

NOROTRIM 24 Inspuiting

Reg. Nr G 2620 (Wet 36/1947)

Slegs vir gebruik deur, of onder die beheer van 'n veearts.

Berging:

Bewaar in 'n koel droë plek. Beskerm teen lig.

Bevat:

Sulfadiazien 200,0 mg en Trimetoprim 40,0 mg per ml.

Aanduidings:

Norotrim 24 is aangewys vir die behandeling van longontsteking, bakteriese diarree, urogenitale infeksies, vrotpootjie, septiese gewrigontsteking en ander infeksies veroorsaak deur sulfonamied-vatbare organismes in perde, beeste, skape, bokke, varke, honde en katte. Die spektrum dek beide Gram-negatiewe en Gram-positiewe bakterieë insluitende: *Actinobacillus spp.*, *Actinomyces bovis*, *Bordetella spp.*, *Corynebacterium spp.*, *Escherichia coli*, *Fusiformis spp.*, *Haemophilus spp.*, *Klebsiella spp.*, *Listeria monocytogenes*, *Nocardia spp.*, *Pasteurella spp.*, *Proteus spp.*, *Salmonella spp.*, *Staphylococcus spp.* en *Streptococcus spp.*

Norotrim 24 word ook aangewys vir ander respiratoriese infeksies van bakteriese oorsprong soos rinitis en bronritis asook ander spysverteringskanaal infeksies.

Waarskuwings:

Melk moenie vir menslike verbruik gebruik word binne 48 uur na die laaste behandeling nie. Beeste moenie vir menslike gebruik geslag word binne 12 dae na die laaste behandeling nie. Varke moenie vir menslike gebruik geslag word binne 15 dae na die laaste behandeling nie. Skape moenie vir menslike gebruik geslag word binne 28 dae na die laaste behandeling nie.

Moenie in diere wat sensief is vir sulfonamide, wat erg lewerbeskadiging toon of wat bloedabnormaliteite wys, gebruik nie. Moenie in perde wat medisyne-verwante hart-aritmie wys, wat gepaardgaan met sekere verdowings- of sedearmiedmiddels gebruik nie.

Moenie inspuitingsroetes gebruik wat nie aanbeveel word nie. Moenie intraperitonalaal inspuit nie. Binneaarse inspuitings moet stadig toegedien word om anafilaktiese skok te vermy. Maak seker dat daar toegang tot genoegsame drinkwater tydens behandlingsperiode is.

Hou buite bereik van kinders, oningepte persone en diere.

Alhoewel hierdie middel breedvoerig onder 'n wye verskeidenheid van toestande getoets is, mag dit faal as gevolg van verskeie redes. Indien dit vermoed word, raadpleeg 'n veearts en verwittig die registrasiehouer.

Voorsorgmaatreëls:

Hierdie produk mag teen lae temperatuure kristalliseer. Die kristalle kan opgelos word deur liggies te verwarm.

Gebruiksaanwysings:

Gebruik slegs soos aangedui.

Perde, beeste, skape, bokke en varke:

1 ml per 16 kg liggaamsmassa daagliks deur middel van binnespiele of stadige binneaarse inspuiting. In gevalle van erge infeksies mag die dosis tot 1 ml per 10 kg liggaamsmassa verhoog word.

Honde en katte:

1 ml per 8 kg liggaamsmassa daagliks deur middel van onderhuidse inspuiting. Die aanbevole toedieningsplek in honde is onder die los vel agter die nek.

'n Enkele inspuiting mag in ongekompliseerde gevalle genoeg wees, maar in erger gevalle mag die dosis vir tot 5 dae herhaal word.

Registrasiehouer:

Biotech Laboratories (Edms) Bpk
Grond Vloer, Blok K Wes,
Central Park, 400 16de Straat,
Midrand, Gauteng, 1685

Remark deur:

Biotech Laboratories (Edms) Bpk
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