FOR ANIMAL USE ONLY

Baycox® 5%

Reg. No. G3662 (Act 36/1947)
Namibia: S0 Reg. No. V07/17.4.1/461
Act 13/2003

NAFDAC Reg No. A10-0271 **POM**

Coccidiocide

Oral suspension

CAUTION

Store in a dry place below 30 °C

Contains per mℓ: **Toltrazuril** 50 mg

For the control of coccidiosis in neonatal piglets.
For the treatment of coccidiosis in cattle and sheep.

For cattle, sheep and piglets

Warnings:
Piglets: Allow 77 days between treatment and slaughter for human consumption.
Cattle: Withdrawal period for meat and offal is 63 days. Do not use in lactating cows producing milk for human consumption. Do not use in pregnant animals.
Sheep: Withdrawal period for meat and offal is 42 days. Do not use in lactating sheep producing milk for human consumption. Do not use in pregnant animals.

Following withdrawal of first dose, use product within 6 months. Discard any unused material.
Keep out of reach of children, uninformed persons and animals. Store away from food and feed.
Although this remedy has been extensively tested under a large variety of conditions, failure may ensue as a result of a wide range of reasons. If this is suspected, seek veterinary advice and notify the registration holder.
Dispose of contents, contaminated packaging, and empty containers to an approved waste disposal plant.

Precautions:
Wash any splashes from skin and eyes immediately with water. Product must be used within 6 months of first use or opening.

Directions for use:
Use only as directed. Shake well before use.

Pigs: Each piglet to be treated on day 3-5 of life with a single oral dose of 20 mg toltrazuril/kg body mass corresponding to 0,4 mℓ oral suspension per kg body mass. Due to the small volumes required to treat individual piglets, use of dosing equipment with a dose accuracy of 0,1 mℓ is recommended. Treatment during an outbreak will be of limited value for the individual piglet because of damage to the small intestine having already occurred.
Cattle: 3 mℓ oral suspension per 10 kg body weight. Each animal should be treated with a single oral dose of 15 mg toltrazuril/kg body weight corresponding to 3mℓ oral suspension per 10 kg body weight.
Sheep: 0.4 mℓ oral suspension per kg body weight. Each animal should be treated with a single oral dose of 20 mg toltrazuril/kg body weight, corresponding to 0.4 mℓ oral suspension per kg body weight.
Cattle and Sheep: To obtain maximum benefit, animals should be treated before the expected onset of clinical signs, i.e. in the prepatent period. This is best achieved in situations where the pattern of the appearance of clinical signs in young or recently introduced animals is well-known and predictable.

Presentation:
100 mℓ, 250 mℓ, 1ℓ, 2ℓ and 5ℓ plastic bottles.

Registration Holder / Registrasiehouer:
Bayer (Pty) Ltd, Animal Health Division
Reg. No.: 1968/011192/07
27 Wrench Rd, Isando, 1601, R.S.A
Tel: (011) 921 5736, Fax: (011) 921 5751
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Bayer

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SLEGS VIR DIEREGEBRUIK

Baycox® 5%

Reg. Nr. G3662 (Wet 36/1947)
Namibië: S0 Reg. Nr. V07/17.4.1/461
Wet 13/2003

NAFDAC Reg No. A10-0271 **POM**

Koksidiosied

Orale suspensie

VERSIGTIG

Berg in 'n droë plek benede 30 °C

Bevat per mℓ: **Toltrazuril** 50 mg

Vir die beheer van koksidiose in pasgebore varkies.
Vir die behandeling van koksidiose in beeste en skape.

Vir beeste, skape en varkies

Waarskuwings:
Varke: Laat 77 dae toe tussen behandeling en slagting vir menslike gebruik.
Beeste: Die onttrekkingsperiode vir vleis en afval is 63 dae. Moenie gebruik in lakterende koeie wat melk vir menslike gebruik produseer nie. Moenie in dragtige diere gebruik nie.
Skape: Die onttrekkingsperiode vir vleis en afval is 42 dae. Moenie gebruik in lakterende skape wat melk vir menslike gebruik produseer nie. Moenie in dragtige diere gebruik nie.

Gebruik die produk binne 6 maande na die ontrekking van die eerste dosis. Vernietig enige ongebruikte produk daarna.

Hou buite bereik van kinders, oningeligte persone en diere. Stoor eenkant, weg van voedsel en voer.

Alhoewel hierdie middel 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. Gooi inhoud, gekontamineerde verpakkingsmateriaal en leë houers weg deur 'n goedgekeurde afval maatskappy.

Voorsorgmaatreëls:
Was enige spatsels dadelik af van die vel en oë met water. Produk moet binne 6 maande gebruik word na oopmaak.

Aanwysings vir gebruik:
Gebruik slegs soos aangedui. Skud goed voor gebruik.

Varkies: Elke varkie moet per bek behandel word op lewensdag 3-5 met 'n enkele 20 mg-dosis toltrazuril/kg liggaamsmassa, ooreenstemmend met 0,4 mℓ orale suspensie per kg liggaamsmassa. As gevolg van die klein volumes wat benodig word om individuele varkies te behandel, word die gebruik van doseringstoerusting met 'n dosis-akkuraatheid van 0,1 mℓ aanbeveel. Behandeling gedurende 'n uitbraak sal van beperkte waarde wees vir die individuele varkie as gevolg van skade aan derms wat alreeds plaasgevind het.
Beeste: 3 mℓ orale suspensie per 10 kg liggaamsgewig. Elke dier moet per bek behandel word met 'n enkele dosis van 15 mg toltrazuril/kg liggaamsgewig, wat ooreenkom met 3 mℓ orale suspensie per 10 kg liggaamsgewig.
Skape: 0.4 mℓ orale suspensie per kg liggaamsmassa. Elke dier moet per bek behandel word met 'n enkele dosis van 20 mg toltrazuril/kg liggaamsmassa, wat ooreenkom met 0,4 mℓ orale suspensie per kg liggaamsgewig.
Beeste en Skape: Om die maksimum voordeel te trek, moet diere behandel word voor die verwagte aanvang van kliniese tekens, d.i. in die prepatente periode. Dit word die beste bereik in situasies waar die voorkomspatroon van kliniese tekens by jong of onlangs ingeslote diere goed bekend en voorspelbaar is.

Aanbieding:
100 mℓ 250 mℓ, 1ℓ, 2ℓ en 5ℓ plastiek bottels.

Manufacturer:
KVP Pharma + Veterinar Produkte GmbH
Projensdorfer Str. 324, D-24106 Kiel
Germany

® is die geregistreerde handelsmerk van Bayer AG, Duitsland

L.ZA.05.2014.0123

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PM no.: 86566591
CR no.: 201806444
Format: 148 x 210 mm
Font size: 6 pt
Drawing no.:

Colours:
Schwarz
Kontur

Operator ID: MK
Version no.: 4
Date: 13.12.2018

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