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### Chanaverm 7.5% Oral Solution

**Species:** Sheep, Cattle

**Therapeutic indication:** **Pharmaceuticals: Endoparasiticides:** Anthelmintics for cattle, Anthelmintics for sheep

**Active ingredient:** Levamisole Hydrochloride

**Product:** Chanaverm 7.5% Oral Solution

**Product index:** Chanaverm 7.5% Oral Solution

**Cattle - milk:** See notes

**Cattle - meat:** 20 days

**Sheep - meat:** 20 days

**Withdrawal notes:** Not for use in animals producing milk for human consumption.

**Incorporating:**

### Presentation

A yellow coloured clear solution for oral administration. Each ml contains 75 mg of Levamisole Hydrochloride. Also contains sodium metabisulphite 0.1% w/v as preservative and Tartrazine (E102) 0.00375% w/v as colourant.

### Uses

Chanaverm 7.5% is a broad spectrum anthelmintic for the treatment and control of gastro-intestinal and pulmonary nematode infections in cattle and sheep. Chanaverm 7.5% is highly effective against mature and developing immature stages of levamisole susceptible major stomach and bowel worm species including, Gastro-intestinal worms: *Haemonchus* spp; *Ostertagia* spp (except inhibited *Ostertagia* larvae in cattle); *Nematodirus* spp; *Trichostrongylus* spp; *Cooperia* spp; *Oesophagostomum* spp; *Chabertia* spp; *Bunostomum* spp; Lungworms: *Dictyocaulus* spp.

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Chanaverm 7.5% is not effective against Type II Winter Scour.

### **Dosage and administration**

Chanaverm 7.5% should be administered as an oral drench. Dosing should be carried out accurately, preferably using a gun system at a rate of 7.5 mg Levamisole hydrochloride per kg bodyweight.

Cattle: 1 ml Chanaverm 7.5% per 10 kg bodyweight.

Sheep: 0.5 ml Chanaverm 7.5% per 5 kg bodyweight.

#### *Dosage guide*

##### **Cattle – Liveweight**

50 kg (approx. 1 cwt)

100 kg (approx. 2 cwt)

150 kg (approx. 3 cwt)

200 kg (approx. 4 cwt)

250 kg (approx. 5 cwt)

300 kg (approx. 6 cwt)

Cattle over 300 kg should be given a further 1 ml for each additional 10 kg bodyweight.

##### **Sheep – Liveweight**

10 kg (approx. 22 lb)

20 kg (approx. 44 lb)

30 kg (approx. 66 lb)

40 kg (approx. 88 lb)

50 kg (approx. 110 lb)

60 kg (approx. 132 lb)

Sheep over 60 kg should be given a further 0.5 ml for each additional 5 kg bodyweight.

### **Contra-indications, warnings, etc**

At normal therapeutic dosages side effects are rarely seen. Overdosage may occasionally result in the appearance of cholinergic type symptoms such as salivation, muscular tremors and head shaking. They are more likely to be observed in cattle than in sheep.

Care should be taken to estimate accurately the bodyweight of animals to be treated before calculating the dosage. Cattle must not be treated within a period of 14 days before or after treatment with organophosphorus compounds or diethylcarbamazine citrate.

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The product may be given to young, pregnant and lactating animals, but due regard must always be paid to the animal's physical condition and the presence of inter-current diseases.

Veterinary advice should be sought on appropriate dosing programmes and stock management to achieve adequate parasite control, and to reduce the likelihood of anthelmintic resistance developing and also if the product does not achieve the desired clinical effect, since other diseases, nutritional disturbances or anthelmintic resistance may be involved. When a dosing gun is used to administer the product, care should be taken to avoid occurrences of dosing gun pharyngitis.

### **Withdrawal periods**

Animals must not be slaughtered for human consumption during treatment. Cattle and sheep may be slaughtered for human consumption only after 20 days from the last treatment. Not for use in animals producing milk for human consumption.

### **Operator warnings**

When using do not eat, drink or smoke. Wash splashes from eyes and skin immediately. If irritation persists consult your doctor. Take off immediately any contaminated clothing. Wash hands and exposed skin before meals and after work. Levamisole can cause idiosyncratic reactions and serious blood disorders in a very small number of people. If symptoms such as dizziness, nausea, vomiting or abdominal discomfort are experienced when using this product, or sore mouth/throat or fever occur shortly afterwards, then medical advice should be sought immediately.

### **Pharmaceutical precautions**

Do not store above 25°C. Protect from light. Do not mix with other products.

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

### **Legal category**

**Legal category:**POM-VPS

### **Packaging quantities**

1 L, 2.5 L and 5 L

### **Marketing Authorisation Number**

Vm 11990/4002.

### **Significant changes**