PARTICULARS TO APPEAR ON THE OUTER PACKAGE

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Chanaverm 7.5% Oral Solution

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Oral solution containing 7.5% w/v Levamisole HCL

3. PHARMACEUTICAL FORM

Oral Solution

4. PACKAGE SIZE

2.5 L
5 L

5. TARGET SPECIES

Cattle and Sheep

6. INDICATION(S)

Chanaverm 7.5% is a broad spectrum anthelminitic for the treatment and control of gastro-intestinal and pulmonary nematode infections in cattle and sheep. Chanaverm 7.5% is effective against mature and developing immature stages of levamisole susceptible major worm species including, Gastro-intestinal worms: *Haemonchus* spp; *Ostertagia* spp (except inhibited *Ostertagia* larvae in cattle); *Nematodirus* spp; *Trichostrongylus* spp; *Cooperia* spp; *Oesophagostomum* spp; *Chavertia* spp; *Bunostomum* spp. Lungworms: *Dictoycaulus* spp. Chanaverm 7.5% is not effective against type II winter scour.

7. METHOD AND ROUTE(S) OF 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 7.5mg Levamisole hydrochloride per kg bodyweight.

To ensure administration of a correct dose, body weight should be determined as accurately as possible; accuracy of the dosing device should be checked.

Cattle: 1ml Chanaverm 7.5% per 10kg bodyweight

Sheep: 0.5ml Chanaverm 7.5% per 5kg bodyweight

Cattle – Liveweight	Dose	Sheep- Liveweight	Dose
50 kg (approx. 1 cwt)	5 ml	10 kg (approx. 22 lb)	1 ml
100 kg (approx. 2 cwt)	10 ml	20 kg (approx. 44 lb)	2 ml
150 kg (approx. 3 cwt)	15 ml	30 kg (approx. 66 lb)	3 ml
200 kg (approx. 4 cwt)	20 ml	40 kg (approx. 88 lb)	4 ml
250 kg (approx. 5 cwt)	25 ml	50 kg (approx. 110 lb)	5 ml
300 kg (approx. 6 cwt)	30 ml	60 kg (approx. 132 lb)	6 ml

Cattle over 300kg should be given a further 1ml for each additional 10kg bodyweight. Sheep over 60kg should be given a further 0.5ml for each additional 5kg bodyweight.

8. WITHDRAWAL PERIOD

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

9. SPECIAL WARNING(S), IF NECESSARY

At normal therapeutic dosages side effects are rarely seen. Overdosage may occasionally result in the appearance of cholongergic 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.

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.

Do not mix with other products

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.

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.

10. EXPIRY DATE

11. SPECIAL STORAGE CONDITIONS

Do not store above 25°C

Protect from light

12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Dispose of any unused product and empty container in accordance with guidance from your local waste regulation authority.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE *[Distribution category]*

POM-VPS

To be supplied only on veterinary prescription.

For animal treatment only

14. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

Keep out of reach of children

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Chanelle Pharmaceuticals Manufacturing Ltd
Loughrea
Co Galway
H62 FH90
Ireland

Manufactured by:
Chanelle Pharmaceuticals Manufacturing Limited
Loughrea
Co Galway
Ireland

16. MARKETING AUTHORISATION NUMBER(S)

Vm 08749/3025

17. MANUFACTURER'S BATCH NUMBER

Gavin Hall
Approved: 11 September 2025