IVERMECTIN- ivermectin oral paste Vetr LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Vetr Ivermectin (1.87% ivermectin oral paste)

Animal Safety

Not safe or approved for human use. May be used in horses of all ages including mares at any stage of pregnancy. KEEP OUT OF REACH OF CHILDREN.

Directions for Use

One syringe contains sufficient paste to treat one 1250 lb horse at the recommended dose rate (0.2 mg/kg) body weight. Do not underdose. Ensure each animal receives a complete dose based on a current body weight. Underdosing may result in ineffective treatment, and encourage the development of parasite resistance.

Storage

Store at room temperature, 20-25 C (68-77 F)

Active Ingredients

Ivermectin USP, 18.7 mg/g (1.87% w/w)

Other Ingredients

Certified vegetable glycerine, natural preservative, stabilizers and flavoring



Oral Paste | Anthelminit and Boticide Removes woms and bots for Horses | with a single dose Vet Wt. 0.21 oz (6.08 g) Contains Single Dose | 1 Syringe

IVERMECTIN PASTE 1.87%



Oral Paste for Horses Anthelmintic and Boticide Removes worms and bots with a single dose

Net Wt. 0.21 oz (6.08 g) Contains Single Dose | 1 Syringe

IVERMECTIN							
ivermectin oral paste							
Product Informat	tion						
Product Type		OTC ANIMAL DRUG	Item Code	(Sourc	e)	NDC	2:86213-771
Route of Administra	tion	Oral					
Active Ingredient	/Active	Moiety					
.		ent Name		Basis	of Streng	gth	Strength
IVERMECTIN (UNII: 8883YP2R6D) (IVERMECTIN - UNII:8883YP2R6D)IVERMECTIN						-	18.7 mg in 1 g
IVERMECTIN (UNII: 8883	31P2R6D) (I	IVERMECTIN - UNII:88831P2	ROD)				10.7 mg mig
IVERMECTIN (UNII: 8883	31P2R6D) (I	VERMECTIN - UNII:88831P2	KOD)	IVERMECT			10.7 mg mig
		VERMECTIN - UNII:88831P2	KOD)	IVERMEC			10.7 mg m i g
		VERMECTIN - UNII:88831P2	KOD)	IVERMEC			10.7 mg m i g
		Ingredient Name		IVERMEC			Strength
	nts			IVERMEC			
Inactive Ingredier	nts 3C0OX)	Ingredient Name		IVERMEC			
Inactive Ingredier GLYCERIN (UNII: PDC6A3	nts 3C0OX) (UNII: 1VPL	Ingredient Name J26JZZ4)		IVERMEC			
Inactive Ingredier GLYCERIN (UNII: PDC6A3 POTASSIUM SORBATE	nts 3C0OX) (UNII: 1VPU : A2I8C7HIS	Ingredient Name J26JZZ4) ƏT)					
Inactive Ingredier GLYCERIN (UNII: PDC6A3 POTASSIUM SORBATE METHYLPARABEN (UNII:	nts 3C0OX) (UNII: 1VPU : A2I8C7HIS	Ingredient Name J26JZZ4) ƏT)					
Inactive Ingredier GLYCERIN (UNII: PDC6A3 POTASSIUM SORBATE METHYLPARABEN (UNII:	nts 3C0OX) (UNII: 1VPU : A2I8C7HIS : Z8IX2SC1	Ingredient Name J26JZZ4) ƏT)					
Inactive Ingredier GLYCERIN (UNII: PDC6A3 POTASSIUM SORBATE METHYLPARABEN (UNII: PROPYLPARABEN (UNII: PROPYLPARABEN (UNII:	nts 3C0OX) (UNII: 1VPU : A2I8C7HI9 2 Z8IX2SC1	Ingredient Name J26JZZ4) ƏT)			ore		
Inactive Ingredier GLYCERIN (UNII: PDC6A3 POTASSIUM SORBATE METHYLPARABEN (UNII: PROPYLPARABEN (UNII: PROPYLPARABEN (UNII:	nts 3C0OX) (UNII: 1VPU : A2I8C7HI9 2 Z8IX2SC1	Ingredient Name J26JZZ4) 9T) OH)			:ore		
Inactive Ingredier	nts 3C0OX) (UNII: 1VPU : A2I8C7HI9 2 Z8IX2SC1	Ingredient Name J26JZZ4) 9T) OH)		Sc	:ore		

Pa	Packaging						
#	ltem Code	Package Description	Marketing	Start Date	Mark	eting End Date	
1	NDC:86213-771-01	6.1 g in 1 SYRINGE, PLASTIC					
Marketing Information							
	Marketing	Application Number or Mo	onograph	Marketing S	tart	Marketing End	
	Category	Citation		Date		Date	
	approved drug ner			05/07/2025			

Labeler - Vetr LLC (132219984)

Establishment						
Name	Address	ID/FEI	Business Operations			
Vetr LLC		132219984	manufacture, label			

Revised: 5/2025

Vetr LLC