

PRIMEGUARD- omeprazole (37%) 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 PrimeGuard (37% omeprazole oral paste)

Not for use in humans. Keep this and all medications out of the reach of children. In case of ingestion by humans, contact a physician. Do not use in horses intended for human consumption. PrimeGuard is intended for use in healthy horses. If you notice any signs of illness prior to or during the use of this product, consult your veterinarian for appropriate diagnosis and treatment recommendations.

The minimum recommended dosage is 1 mg/kg per day (0.45 mg/lb) or 1/4 syringe. Each syringe contains 4 individual daily doses for horses weighing 600-1200 lbs. Please refer to this dosage chart for help in determining the correct dose for your horse:

Horse WeightDose*

<600 lbs Consult your veterinarian

600-1200 lbs 1 dose per day

>1200 lbs 2 doses, once per day

*Dosage duration: 8 or 28 days. Consult your veterinarian.

Store at 68°F-77°F (20-25°C). For transport, brief periods between 59°F-86°F (15-30°C) are permitted.

Omeprazole USP 2.28g per syringe (37% omeprazole)





PRIMEGUARD

omeprazole (37%) oral paste paste

Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-221
Route of Administration	oral		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9)	OMEPRAZOLE	0.37 g in 1 g

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	APPLE	Imprint Code	

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-221-01	1 in 1 CARTON		
1		6.16 g in 1 SYRINGE, PLASTIC		
2	NDC:86213-221-02	6 in 1 CARTON		
2		6.16 g in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/26/2026	

PRIMEGUARD

omeprazole (37%) oral paste paste

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-222
Route of Administration	oral		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9)	OMEPRAZOLE	0.37 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor	CINNAMON	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-222-01	1 in 1 CARTON		
1		6.16 g in 1 SYRINGE, PLASTIC		
2	NDC:86213-222-02	6 in 1 CARTON		
2		6.16 g in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/26/2026	

PRIMEGUARD

omeprazole (37%) oral paste paste

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86213-223
Route of Administration	oral		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9)	OMEPRAZOLE	0.37 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor	MALT (molasses)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86213-223-01	1 in 1 CARTON		
1		6.16 g in 1 SYRINGE, PLASTIC		
2	NDC:86213-223-02	6 in 1 CARTON		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/26/2026	

Labeler - Vetr, LLC (132219984)**Establishment**

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

Revised: 1/2026

Vetr, LLC