EXOCOUGH- potassium iodide and ammonium chloride solution Nu-Vet Pty Ltd

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.

Active

POTASSIUM IODIDE 20 g / L AMMONIUM CHLORIDE 20 g /L

Inactive ingrdients

Glycerin, H20 Distilled Water, Benzoyl Alcohol and Red Dye

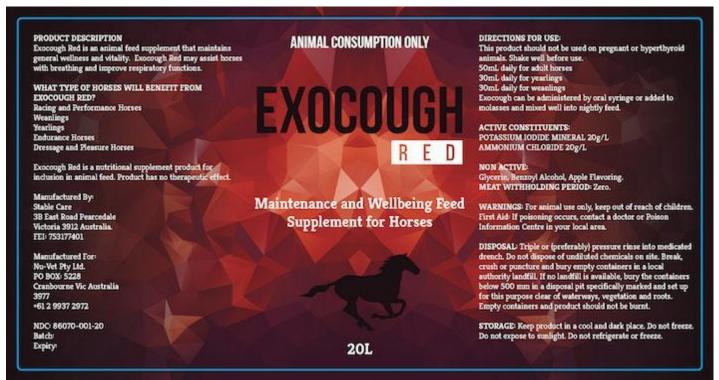
Purpose of use

Equine Health Supplement

Do not use

Not to be used for horses in human consumption.

Product label



EXOCOUGH

potassium iodide and ammonium chloride solution

Product Type		OTC ANIMAL DRUG Item Code (Source)			NDC:86070-001		
Route of Administration	1	ORAL					
Active Ingredient/A	ctive Moi	ety					
Ingredient Name E						f Strength	Strengt
POTASSIUM IODIDE (UNII: 1C4QK22F9J) (IODIDE ION - UNII:09G4I6V86Q)						POTASSIUM IODIDE	
AMMONIUM CHLORIDE (UNII: 01Q9PC255D) (AMMONIUM CATION - UNII:54S6852014) AMMO						IUM CATION	20 g in 1 I
Inactive Ingredients							
		Ingredient Name				Strength	
GLYCERIN (UNII: PDC6A3	BCOOX)						
WATER (UNII: 059QF0KO	0 R)						
BENZYL ALCOHOL (UNI	I: LKG8494V	VBH)					
Product Characteris	tics						
Color red		red	Score				
Shape			Size				
Flavor	L	APPLE	Imprint Code				
Contains							
Packaging							
# Item Code	Pac	kage Description	Marketing Start Date		Marketing End Date		ıd Date
1 NDC:86070-001-20	20 L in 1	DRUM					
I NDC.00070-001-20							
	.•						
Marketing Inform							
		on Number or Monogra	aph Citation	Marketing St 07/07/2016	art Date	Marketing	g End Date

Labeler - Nu-Vet Pty Ltd (750770047)

Establishment									
Name	Address	ID/FEI	Business Operations						
Stable Care Pty		753177401	manufacture						

Establishment

Name	Address	ID/FEI	Business Operations
G AMPHRAY LABORATORIES		862288982	api manufacture