TOTAL ANTIMONY - antimony potassium tartrate liquid White Manufacturing Inc. DBA Micro -West

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts

Active Ingredients

antimonium tartaricum 12x,30x,200c,1m

Uses

Metallic taste, gout-like symptoms, angina, anorexia, fatigue, myopathy, hypotension,

Keep Out of Reach of Children

KEEP OUT OF REACH OF CHILDREN

INDICATIONS

INDICATIONS: Metallic taste, gout-like symptoms, angina, anorexia, fatigue, myopathy, hypotension, "antimony spots" if exposed to vapor

Ask Doctor

If pregnant or nursing, please consult physician before using this product

Directions

Directions: Orally 6 drops, 4 times a day

Inactive Ingredient

Not more than 20% Alcohol

Principal Display Panel



INDICATIONS: Metallic taste, gout-like symptoms, angina, anorexia, fatigue, myopathy, hypotension, "antimony spots" if exposed to vapor.

Lot#



4 Tier Multi-Potency Total Antimony

NDC# 61657-0966-1 1 Fl. oz., 30 ml. Not more than 20% alcohol

INGREDIENTS: ANTIMONIUM TARTARICUM, 12X, 30X, 200C, 1M

WARNING: If pregnant or nursing, please cian before using this product. KEEP OUT OF REACH OF CHILDREN

Manufactured for Micro West Douglas, WY 82633 1-307-358-5066

TOTAL ANTIMONY

antimonium tartaricum liquid

Prod	luct	Info	rma	tion
Proc	luct	11110	miliai	uon

Product Type HUMAN OTC DRUG LABEL Item Code (Source) NDC:61657-0966 Route of Administration ORAL **DEA Schedule**

	Active Ingredient/Active Moiety	
- 1		

Ingredient Name Basis of Strength Strength ANTIMONY POTASSIUM TARTRATE (ANTIMONY CATION (3+)) ANTIMONY POTASSIUM TARTRATE 12 [hp_X] in 30 mL

Inactive Ingredients

Ingredient Name Strength Alcohol

Packaging

ı				
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:61657-0966-1	30 mL in 1 BOTTLE, DROPPER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/13/2011	

Labeler - White Manufacturing Inc. DBA Micro -West (082307831)

Establishment				
Name	Address	ID/FEI	Business Operations	
White Manufacturing Inc. DBA Micro -West		082307831	label(61657-0966), pack(61657-0966)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Washington Homeopathic Products		084929389	manufacture(61657-0966)	

Revised: 9/2012 White Manufacturing Inc. DBA Micro -West