ARSENICUM SULPHURATUM FLAVUM- arsenic trisulfide liquid Washington Homeopathic Products

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

ARSENICUM SULPHURATUM FLAVUM

USES

To relieve the symptoms of itchy skin.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

ARSENICUM SULPHURATUM FLAVUM Itchy skin

STOP USE AND ASK DOCTOR

If symptoms persist/worsen or if pregnant/nursing, consult your practitioner.

DIRECTIONS

Adults: 4 drops into a tsp. of water 3 times a day. Children: 1/2 dose. Repeat at greater intervals as condition subsides. Or as directed bya lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of ARSENICUM SULPHURATUM FLAVUM is 6x–30x, 3c–30c, 200c, 1m, 10m, 50m, and CM.



Availability is subject to change.

All WHP single remedies are made to order; thus, the labels are printed on the same label stock, as the orders are filled.

'Bottle Size,' 'Potency,' and 'Alcohol Percentage' vary on the label depending on customer choice.

Standard bottle sizes for dilution-form remedies are 15ml, 30ml, 50ml, and 100ml.

ARSENICUM SULPHURATUM FLAVUM							
arsenic trisulfide liquid							
Product Information							
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:71919-087			
Route of Administration	ORAL						
Active Ingredient/Active Moi	ety						
Ing	redient Name		Basis of Stre	ngth	Strength		
ARSENIC TRISULFIDE (UNII: 44SIJ80 UNII:C96613F5AV)	0OX) (ARSENIC CATION (3+) -		ARSENIC TRISU	JLFIDE	30 [hp_C] in 1 mL		
Inactive Ingredients							

		Ingredient Name				Strength		
ALCOHOL (UNII: 3K9958V90M)								
WATER (UNII: 059QF0KO0R)								
Product Characteristics								
Color	white (white) Score							
Shape	Size							
Flavor	avor Imprint Code			Code				
Contains								
Packaging								
# Item Code		Package Description		Marketing Date		Marketing Date	-	
1 NDC:71919-087- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product		12/14/2010					
2 NDC:71919-087- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Pro		n Product	12/14/2010				
3 NDC:71919-087- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		ı tio n	12/14/2010				
4 NDC:71919-087- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product			12/14/2010				
Marketing Information								
Marketing Categ	ory Ap	plication Number or Monograph Cit	ation I	Marketing Star	t Date	Marketing En	nd Date	
unapproved homeopathic		12	12/14/2010					

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(71919-087)

Revised: 12/2018

Washington Homeopathic Products