VENOMX - zinc acetate liquid
Phillips Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

DRUG FACTS

Questions and Side effects
Use of this product results in no known side effects when used according to directions. Phillips Company, 311 Chickasaw Street, Millerton, OK USA 74750; Tel. 580-746-2430
Email address: hp@valliant.net

ACTIVE INGREDIENTS

Active Ingredients	Purpose
Zinc acetate (.1% by volume)	Skin Protectant

ASK DOCTOR

Ask a doctor
If symptoms last for more than 7 days or clear up and recur within a few days.
If the conditions do not improve.
Before use on children under 2 years.
Before use if you are allergic to any ingredients listed on this label.

DO NOT USE

Warnings
For external use only
Keep away from children
Avoid contact with eyes
May be harmful if swallowed or inhaled

CHILDREN

Keep away from children
Avoid contact with eyes
May be harmful if swallowed or inhaled

PURPOSE

Active Ingredients	**Purpose**
Zinc acetate (.1% by volume)	Skin Protectant

STOP USE

If symptoms last for more than 7 days or clear up and recur within a few days.
If the conditions do not improve.
Before use on children under 2 years.
Before use if you are allergic to any ingredients listed on this label.

WHEN USING

For external use only
Avoid contact with eyes
May be harmful if swallowed or inhaled
If symptoms last for more than 7 days or clear up and recur within a few days.
If the conditions do not improve.
Store at 40 to 120 degrees F.

WARNINGS

Warnings
For external use only
Keep away from children
Avoid contact with eyes
May be harmful if swallowed or inhaled
Use of this product results in no known side effects when used according to directions.

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DOSAGE & ADMINISTRATION

Uses
Temporarily protects injured or exposed skin or mucous membrane surfaces from harmful or annoying stimuli and may help provide relief to such surfaces

Directions
Apply to affected skin area and rub gently. Repeat the process as needed.

USE

Uses
Temporarily protects injured or exposed skin or mucous membrane surfaces from harmful or annoying stimuli and may help provide relief to such surfaces

Apply to affected skin area and rub gently. Repeat the process as needed.

INACTIVE INGREDIENTS

Inactive ingredients
Water, color, glycerin, hydroxethylcellulose, chlor-hexidine gluconate, glucono delta lactone, methylparaben, sodium hydroxide,sodium dodecylbenzene sulfonate, dimethyl sulfoxide, witch hazel, ascorbic acid, magnesium stearate, silica and stearic acid.

Image of product

Image of product

VENOMX

zinc acetate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43074-207
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
zinc acetate (UNII: FM5526K07A) (zinc - UNII:J41CSQ7QDS)	zinc	0.001 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
dimethyl sulfoxide (UNII: YOW8V9698H)	
ascorbic acid (UNII: PQ6CK8PD0R)	
dipropylene glycol (UNII: E107L85C40)	
water (UNII: 059QF0KO0R)	
glycerin (UNII: PDC6A3C0OX)	
chlorhexidine gluconate (UNII: MOR84MUD8E)	
calcium gluconate (UNII: SQE6VB453K)	
methylparaben (UNII: A2I8C7HI9T)	
sodium hydroxide (UNII: 55X04QC32I)	
sorbic acid (UNII: X045WJ989B)	
magnesium stearate (UNII: 70097M6I30)	
stearic acid (UNII: 4ELV7Z65AP)	
witch hazel (UNII: 101I4J0U34)	
sodium dodecylbenzenesulfonate (UNII: 554127163Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43074-207-02	1 in 1 BOTTLE, PLASTIC		
1	NDC:43074-207-01	3 mL in 1 BOTTLE, DROPPER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	08/23/2010	

Labeler - Phillips Company (612368238)

Establishment

Name	Address	ID/FEI	Business Operations
Phillips Company		612368238	manufacture

Revised: 9/2010 Phillips Company