ADVANCED JELLYFISH STING KIT - zinc acetate powder, for suspension
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

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Uses
Temporarily protects injured or exposed skin or mucous membrane surfaces from harmful or annoying stimuli, and may help provide relief to such surfaces. Contains surfactants to break down oily toxins.

Warnings
n For external use only; do not swallow. n Keep away from children.
n Do not use in the eyes or apply over large areas of the body.
n Do not use if allergic to any ingredient listed on this label.

Directions
Step 1: Wet the skin with water and apply powder to the affected area. Step 2: Use water to wet the special cloth and use it to scrub the affected area to remove jellyfish barbs and to mix the protectant with any remaining toxins. Step 3: Repeat Steps 1 and 2 a minimum of three times or until stinging and pain stops. Step 4: Seek immediate medical attention.

Other information
Store at 40 to 120 degrees F. Net contents of kit: One special scrub cloth and one powder vial. Net contents of powder vial: 3 mL. This product contains no alcohol; no animal products; no biological products.

Inactive ingredients
Water, calcium carbonate, sodium hypochlorite, sodium carbonate, sodium dodecylbenzene sulfonate, ascorbic acid, magnesium stearate, sorbic acid, stearic acid, diproplyene glycol, dimethyl sulfoxide, color.

Questions and Side effects
Report any side effects to Phillips Company, 311 Chickasaw St, Millerton, OK USA 74750
Email: phillips@123phillips.com

ACTIVE INGREDIENTS

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

ASK DOCTOR

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CHILDREN

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PURPOSE

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WHEN USING

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DOSAGE & ADMINISTRATION

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Image of product

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43074-114
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)	
sorbic acid (UNII: X045WJ989B)	
magnesium stearate (UNII: 70097M6I30)	
stearic acid (UNII: 4ELV7Z65AP)	
sodium dodecylbenzenesulfonate (UNII: 554127163Y)	
calcium carbonate (UNII: H0G9379FGK)	
sodium carbonate (UNII: 45P3261C7T)	
sodium hypochlorite (UNII: DY38VHM5OD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:43074-114-18	1 in 1 PACKAGE, COMBINATION		
1	NDC:43074-114-16	3 mL in 1 VIAL, PLASTIC		
2	NDC:43074-114-17	0.003 mL in 1 APPLICATOR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	11/12/2010	

Labeler - Phillips Company (612368238)

Establishment

Name	Address	ID/FEI	Business Operations
Phillips Company		612368238	manufacture