NOSK EX SLEEP- menthol liquid
DreamAir Co.,

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.

ACTIVE INGREDIENT

Active Ingredient: Menthol 2.0%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Alcohol, Glycerin, Dipropylene glycol, Gossypium Herbaceum(cotton) seed extract, Magnolia Kubus Bark extract, Lilium tigrinum extract, Paeonia albiflora root extract, Carbomer, Disodium EDTA, Triethanolamine

PURPOSE

Purpose: Skin Refresher

Caution

Caution: - For the external use only . - Do not apply to eyes ,wounds , broken skin and deep puncture wounds - For 7 years and older only. - When using this product do not get into eyes, Stop use and ask a doctor if rash occurs - Keep out of reach of children - It is recommend to stop applying and to talk to dermatologist immediately if you are suffering such conditions a rash, swell up and itchness - Do not leave it on high, low temperature or exposed on sunlight

KEEP OUT OF REACH OF CHILDREN

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INDICATIONS & USAGE

Indications and usage: 1. Apply to back of neckand, cervical vertebrae and shoulder , make you awake and refreshment. 2. Shake well up and down before use Push a little the ball on apply , liquid easily out.

DOSAGE & ADMINISTRATION

Dosage and administration: Take an adequate amount of this product.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NOSK EX SLEEP

menthol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70767-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Menthol (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	Menthol	0.2 g in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Alcohol (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70767-010-02	6 in 1 CARTON	05/01/2016	
1	NDC:70767-010-01	10 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/01/2016	

Labeler - DreamAir Co., (689846965)

Registrant - DreamAir Co., (689846965)

Establishment

Name	Address	ID/FEI	Business Operations
DreamAir Co.,		689846965	manufacture(70767-010)

Revised: 6/2016 DreamAir Co.,