WET WIPES- salicylic acid swab HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD.

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.

61312004

Active ingredient

salicylic acid 0.5%

Purpose

acne treatment

Use

for the treatment of acne

Warnings

for external use only when using this product

with other topical acne medications ,at the same time or immediately afterwards ,increased skin dryness ,redness or

irritation may occur .If this happens ,only one medication should be used unless directed by a doctor. avoid contact with eyes .if contact occurs, rinse thoroughly with water. keep away from mouth and lips. stop use and ask doctor if skin irritation occurs.

Keep out of reach of children, If swallowed, get medical help or contact a Poison Control Center immediately.

Keep out of reach of children

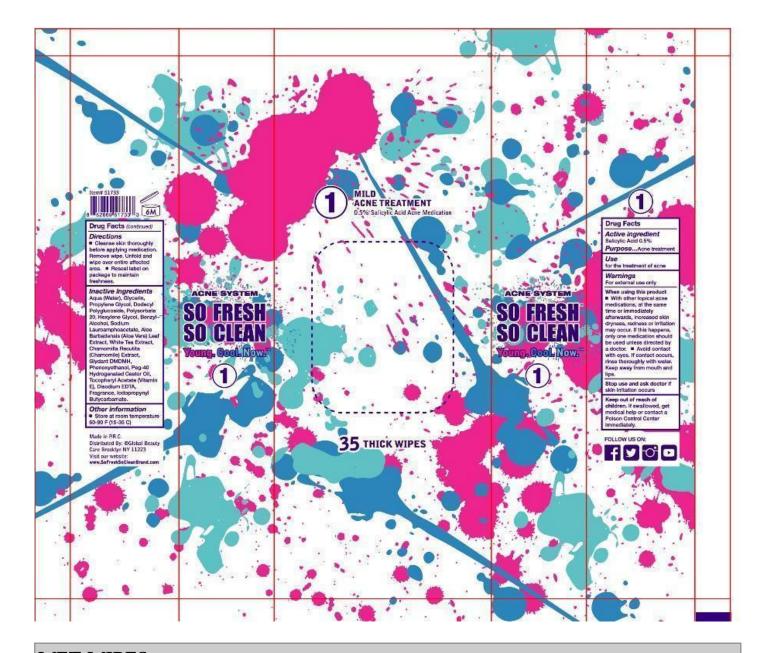
Keep out of reach of children, If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

cleanse skin thoroughly before applying medication ,remove wipe ,unfold and wipe over entire affected area.

reseal label on package to maintain freshness.

aqua (water),glycerin, propylene glycol , dodecyl polyglucoside, polysorbate 20,HEXYLENE GLYCOL , BENZYL ALCOHOL ,SODIUM LAURAMINOPROPIONATE ,ALOE BARBADENSIS(ALOE VERA)EXTRACT ,WHITE TEA EXTRACT ,CHAMOMILLA RECUTITA(CHAMOMILE)EXTRACT ,GLYDANT DMDM H,PHENOXYETHANOL ,PEG-40 HYDROGENATED CASTOR OIL,TOCOPHERYL ACETATE(VITAMIN E) ,DISODIUM EDTA,IODOPROPYNYL BUTYLCARBAMATE



WET WIPES

salicylic acid swab

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61312-004
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.5 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)	
DODECYL PROPIONATE (UNII: 7G537BMD9L)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
HEXYLENE GLYCOL (UNII: KEH0 A3F75J)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM LAURAMINO PRO PIO NATE (UNII: X5NJA9 HXPU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TEA LEAF OIL (UNII: VC855RRT77)	
CHAMO MILE FLO WER O IL (UNII: 60 F80 Z61A9)	
DMDM HYDANTO IN (UNII: BYR0546TOW)	
PHENO XYETHANOL (UNII: HIE492ZZ3T)	
PEG-40 HYDRO GENATED LANOLIN (UNII: 959 P5CXE48)	
TO COPHERYL RETINO ATE (UNII: 0 WN694NBMM)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging					
# Item Co	le	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:61312-00	04-01 35 in 1 B.	AG	0 1/0 1/20 18		
1	0.00054	g in 1 POUCH; Type 0: Not a Combination Product			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part358B	02/25/2015			

Labeler - Hangzhou Guoguang Touring Commodity Co., Ltd. (526890634)

Establishment					
Name	Address	ID/FEI	Business Operations		
HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD.		526890634	manufacture(61312-004)		

Revised: 1/2018

HANGZHOU GUOGUANG TOURING COMMODITY CO., LTD.