

ACNE WASH KISSABLE- salicylic acid gel BRANDS INTERNATIONAL

Kissable Acne Wash Oil Free

Active Ingredient Purpose

Salicylic Acid 2%Acne Treatment

for the treatment of Acne

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

For treatment of acne

Clean twice a day

Wet face, apply to hands add water and work into a lather

Massage fae gently

Rinse thoroughly

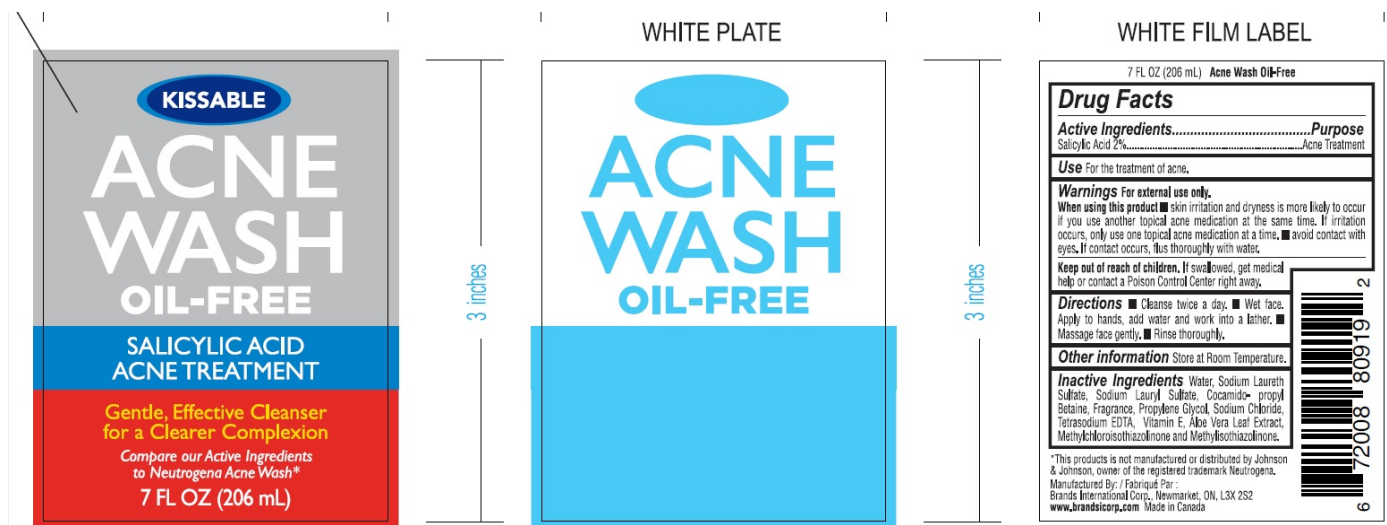
Warning: for external use only.

Ask a doctor or pharmacist before use if you are using other topical acne medications at the same time or immediately following use of this product. This may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

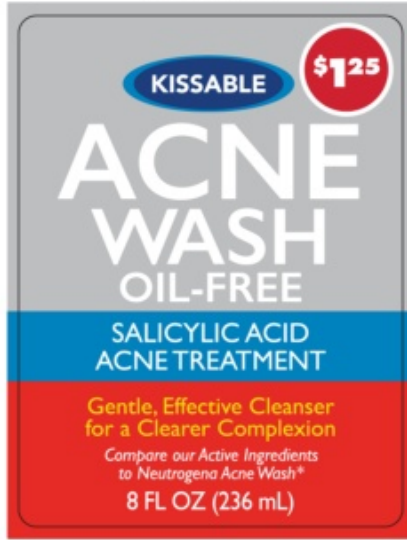
When using this product avoid contact with eyes. If contact occurs, flush thoroughly with water.

Store at room temparture

Inactive Ingredients:water, cocamidopropyl betaine, propylene glycol, sodium chloride, fragrance, Tetra sodium EDTA, aloë barbadensis leaf extract, methylchlorisothiazolinone, methylisothiazoline,



CLEAR FILM LABEL
3.0 X 2.25 in.



WHITE FILM LABEL
3.0 X 2.25 in.



- PMS 287
- PMS Process Blue
- PMS 485
- PMS Yellow C
- Process Black
- White RS

ACNE WASH KISSABLE

salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50157-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3011KX)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-012-20	206 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2023	
2	NDC:50157-012-23	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/27/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/27/2023	

Labeler - BRANDS INTERNATIONAL (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International		243748238	manufacture(50157-012)

Revised: 4/2024

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