BANSBAO SKIN BRIGHTENING AND SPOTS-REMOVING- salicylic acid cream
Mengfei Li Inc.

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.

BANSBAO SKIN BRIGHTENING & SPOTS-REMOVING CREAM

Drug Facts

Active ingredients

Salicylic Acid 2%

Purpose

Acne treatment

Uses

For brightening, removing acne, and spots.

Warnings

- **For external use only.**
- **When using this product**. skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time, if irritation occurs, use only one topical acne medication at a time, Avoid contact with eyes. If contact occurs, flush thoroughly with water.

Keep out of reach of children. If swallowed, get medical help or contact poison control center right away.

Direction. Cleanse twice a day

- Wet face. Apply to hands, add water and work into a lather.
- Massage face gently.

Other Information:

Store at room temperature

Inactive ingredients:

Water, glycerin, propylene glycol, squalane, dimethicone, cyclomethicone, hydroxyethyl urea, glycine, ammonium polyacryloyldimethyl taurate, tocopherol, glucose, mannose, polyglucuronic acid, sodium hyaluronate, Prunus persica (peach) flower extract, Nelumbo nucifera flower extract, Hibiscus militaris flower extract, methylparaben, caprylhydroxamic acid, glyceryl caprylate, disodium EDTA, fragrance.

Bansbao

PRO-CARE

A medicated cream for brightening, removing acne, and spots.

Made in USA

U.S. Distributor:

Mengfei Li Inc. DBA Bansbao USA

Arcadia, CA 91007

www.bansbaousa.com

Packaging

DRUG FACTS

NDC number - 72889-526-95

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BANSBAO SKIN BRIGHTENING AND SPOTS-REMOVING

salicylic acid cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72889-526
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 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SQUALANE (UNII: GW89575KF9)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CYCLOMETHICONE (UNII: NMQ347994Z)	
HYDROXYETHYL UREA (UNII: MBQ7DDQ7AR)	
GLYCINE (UNII: TE7660XO1C)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (55000 MPA.S) (UNII: F01RIY4371)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ANHYDROUS DEXTROSE (UNII: 5SL0G7R0OK)	
MANNOSE, L- (UNII: 2W3YE50TX8)	
POLYGALACTURONIC ACID (UNII: VV3XD4CL04)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
PRUNUS PERSICA FLOWER (UNII: 19GWB0JENH)	
NELUMBO NUCIFERA FLOWER (UNII: 61W322NLDV)	
HIBISCUS LAEVIS FLOWER (UNII: 10813RME15)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
GLYCERYL MONOCAPRYLATE (UNII: TM2TZD4G4A)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
FAGRAEA BERTEROANA FLOWER OIL (UNII: 6665W245VP)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72889-526-95	1 in 1 BOX	01/20/2019	
1		50 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/20/2019	

Labeler - Mengfei Li Inc. (116958347)

Revised: 4/2019 Mengfei Li Inc.