ISLEAF SATIN LIP AND CHEEK DUO CORAL ORANGE- dimethicone liquid
C3 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.

Drug Facts

dimethicone

wter butylene glycol, beeswax, etc

for a gradient effect

keep out of reach of the children

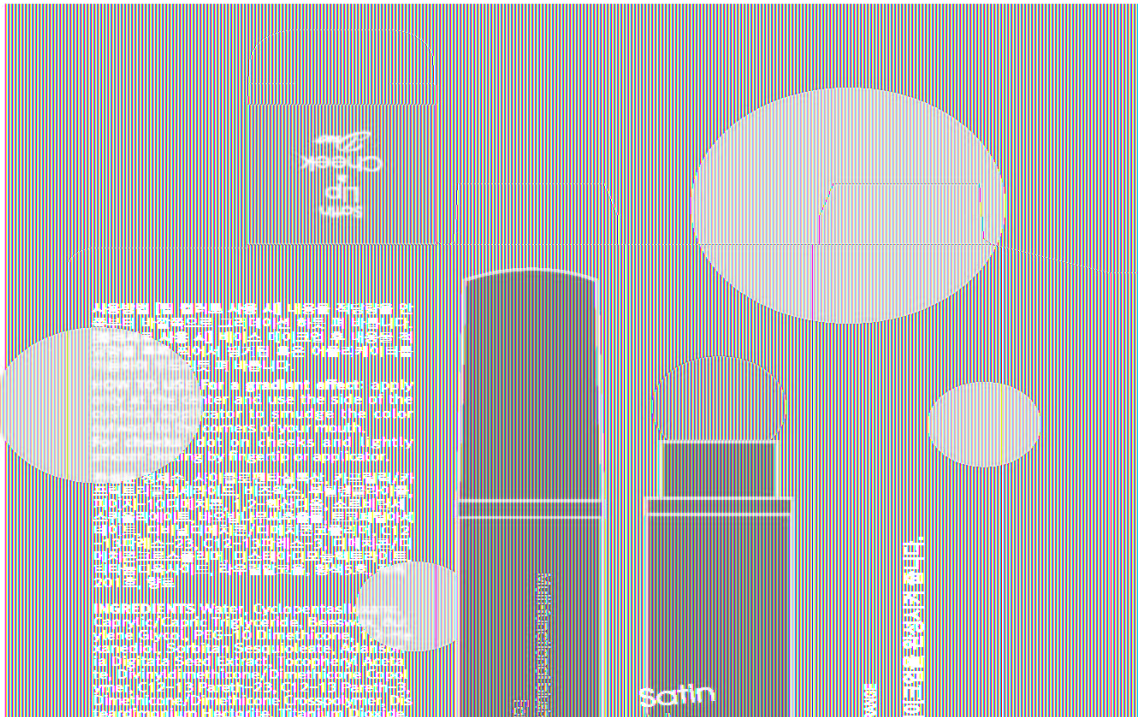
For a gradient effect, apply only at the center and use the side of the cushion applicator to smudge the color outward to the corners of your mouth.

For cheeks, dot on cheeks and lightly smooth dotting by fingertip or applicator

1. If the following symptoms occur after product use, stop using the product immediately and consult a dermatologist (continuous use can exacerbate the symptoms).
1) Occurrence of red spots, swelling, itchiness, and other skin irritation
2) If the symptoms above occur after the application area is exposed to direct sunlight
2. Do not use on open wounds, eczema, and other skin irritations
3. Precaution for Storage and Handling
1) Close the lid after use
2) Keep out of reach of infants and children
3) Do not to store in a place with high/low temperature and exposed to direct sunlight

4. Use as avoiding eye areas.

for external use only



Satin Lip & Cheek Duo CORAL ORANGE

ISLEAF SATIN LIP AND CHEEK DUO CORAL ORANGE

dimethicone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70818-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	0.15 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70818-005-01	4.5 g in 1 APPLICATOR; Type 0: Not a Combination Product	08/08/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	08/08/2017	

Labeler - C3 Co., Ltd. (689846633)

Registrant - C3 Co., Ltd. (689846633)

Establishment

Name	Address	ID/FEI	Business Operations
C3 Co., Ltd.		689846633	label(70818-005) , manufacture(70818-005) , pack(70818-005)

Revised: 8/2017 C3 Co., Ltd.