

ISLEAF CUSHION CONCEALER 23- titanium dioxide 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

titanium dioxide

wter butylene glycol, beeswas, etc.

brighten around eyes or face

keep out of reach of the children

Click end to release product into brush, then apply to areas you want to perfect and brighten around eyes or face. Repeat until desired coverage is achieved, then pat into skin with fingertips to blend flawlessly.

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

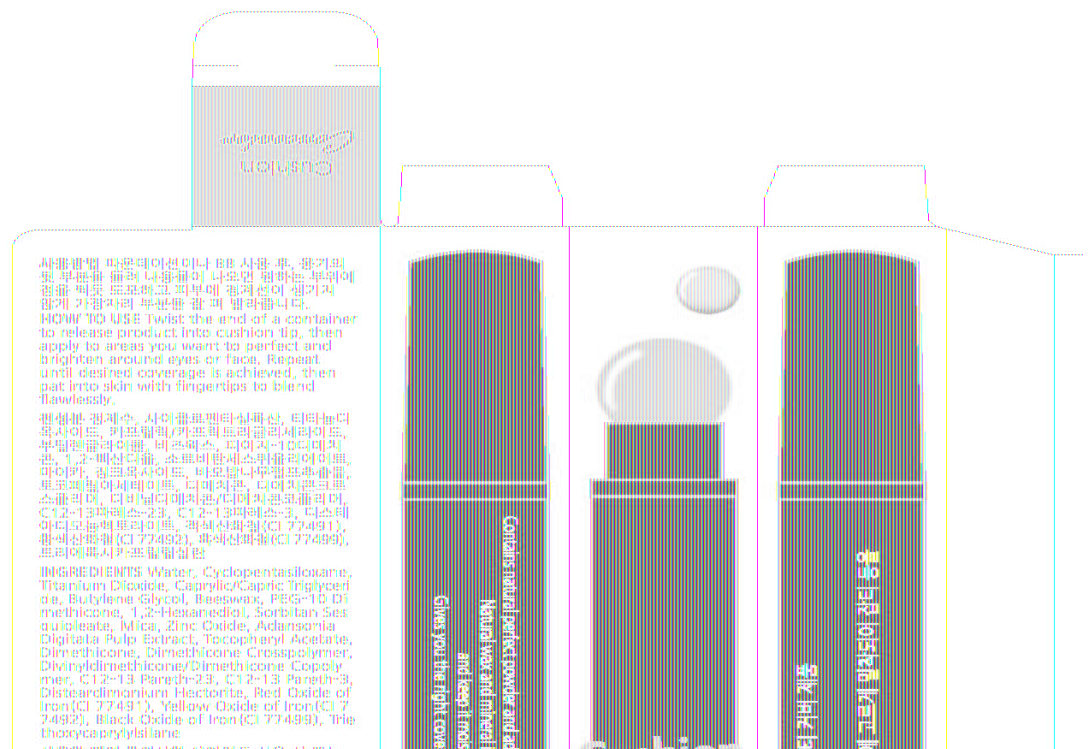
2. Do not use on open wounds, eczema, and other skin irritations

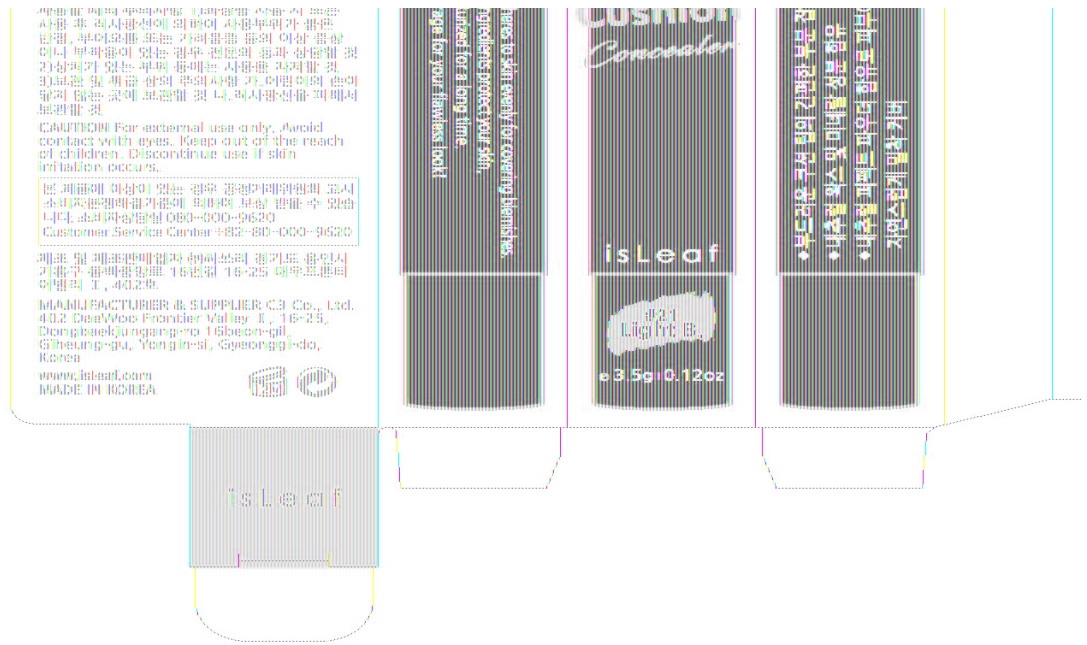
3. Precaution for Storage and Handling

- 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





ISLEAF CUSHION CONCEALER 23

titanium dioxide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70818-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	1.6 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALLANTOIN (UNII: 344S277G0Z)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70818-002-01	3.5 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(70818-002) , manufacture(70818-002) , pack(70818-002)

Revised: 8/2017

C3 Co., Ltd.