ISLEAF SHINE SMOOTH LIP GLAZE PEARL RED- mineral oil 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

mineral oil

polybutene, lauryl alcohol, etc

skin protectant

keep out of reach of the children

Open the cap, press the button to put suitable amount on the applicator and apply evenly. Keep in a place away from direct sunlight with the cap closed after use

- 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 SHINE SMOOTH LIP GLAZE PEARL RED

mineral oil liquid

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:70818-013 |
| Route of Administration | TOPICAL | | |

| Active Ingredient/Active Moiety | | | |
|--|-------------------|---------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP) | MINERAL OIL | 8 g in 100 mL | |

| Inactive Ingredients | | |
|---|---------------|--|
| Ingredient Name | Strength | |
| SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W) | 2 g in 100 mL | |

| Packaging | | | |
|------------------------|---|-------------------------|-----------------------|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 NDC:70818-013- 01 | 2.5 mL 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(70818-013), pack(70818-013), manufacture(70818-013) |

Revised: 8/2017 C3 Co., Ltd.