BDR RE-ACTION DEEP LOW BASE SKIN REFINER EXFOLIATION ESSENCE- glycolic acid liquid

Hanaim International Llc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

bdr Re-action deep low base skin refiner exfoliation essence

Active Ingredients

Glycolic Acid (10%)

Purpose

Anti-wrinkle

Keep out of reach of children

Do not swallow. In case of accidental ingestion, get medical help or contact a Poison Control Center right away.

Warnings

For external use only

Do not use when Your skin is red, inflamed, irritated or painful

When using this product

Do not apply on other parts of the body

Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

☐ Do not apply directly to wound or open cut.

Stop use and ask doctor if rash or irritation on skin develops and lasts.

Keep out of reach of children

Do not swallow. In case of accidental ingestion, seek professional assistance.

Store at room temperature

Uses

Helps to remove dead skin cells gently, and improve skin texture to look radiant and fresh. For large-pored, oily, sensitive skin.

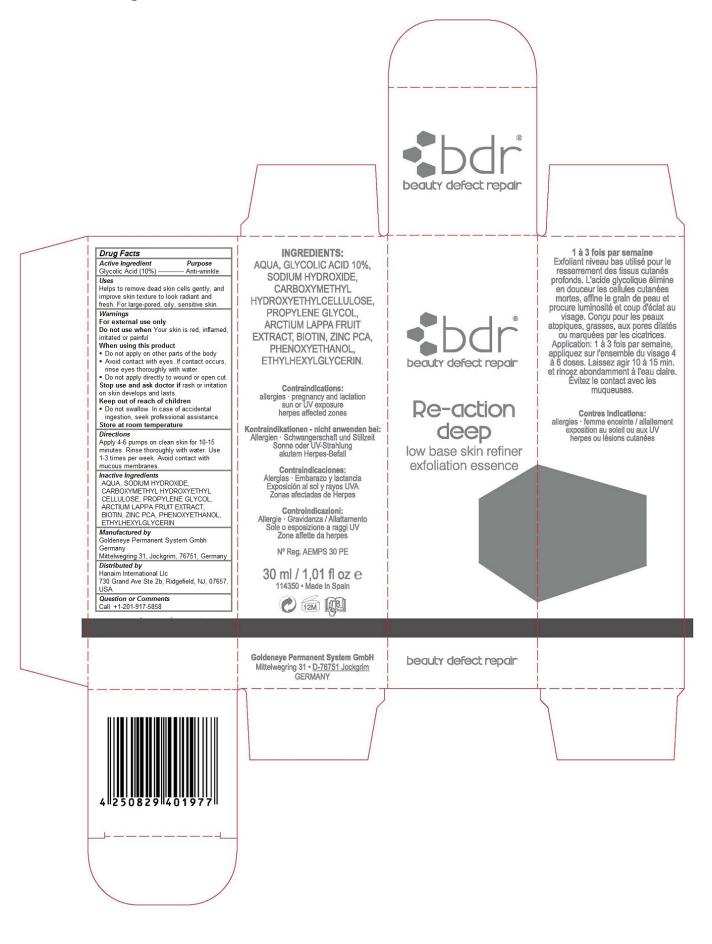
Directions

Apply 4-6 pumps on clean skin for 10-15 minutes. Rinse thoroughly with water. Use 1-3 times per week. Avoid contact with mucous membranes.

Inactive Ingredients

AQUA, SODIUM HYDROXIDE, CARBOXYMETHYL HYDROXYETHYL CELLULOSE, PROPYLENE GLYCOL, ARCTIUM LAPPA FRUIT EXTRACT, BIOTIN, ZINC PCA, PHENOXYETHANOL, ETHYLHEXYLGLYCERIN

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glycolic acid liquid

| Droduct | Information |
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| PIONICI | 111101 1111411011 |

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71056-103

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength | | |
|--|-------------------|--------------|--|--|
| GLYCOLIC ACID (UNII: 0 WT12SX38S) (GLYCOLIC ACID - UNII:0 WT12SX38S) | GLYCOLIC ACID | 3 g in 30 mL | | |

Ingredient Name Strength WATER (UNII: 059QF0KO0R)

SODIUM HYDROXIDE (UNII: 55X04QC32I) **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3)

ARCTIUM LAPPA FRUIT (UNII: EA541308MV)

BIOTIN (UNII: 6 S O 6 U 10 H 0 4)

Inactive Ingredients

ZINC PIDOLATE (UNII: C32PQ86DH4)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)

Packaging # Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:71056-103-02 1 in 1 PACKAGE 11/14/2016

1 NDC:71056-103-01 \mid 30 mL in 1 BOTTLE; Type 0: Not a Combination Product

Marketing Information

| The first matter | | | | |
|-----------------------|--|----------------------|--------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| unapproved drug other | | 11/14/2016 | | |
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$\boldsymbol{Labeler} \textbf{ -} \textbf{ Hanaim International Llc (070319425)}$

$\pmb{Registrant - \text{Hanaim International Llc } (070319425)}$

| Establishment | | | | | | | |
|--------------------------|---------|-----------|---------------------|--|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | | |
| Hanaim International Llc | | 070319425 | relabel(71056-103) | | | | |

Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---------|-------------|------------------------|
| Goldeneye Permanent System Gmbh Germany | | 329 178 144 | manufacture(71056-103) |

Revised: 2/2019 Hanaim International Llc