RUBY REBORN TRANSFUSION DNA KIT- glycerin liquid
Karatica 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.

Glycerin

Adenosine, Ruby, Diamond, DNA, etc

Anti-Wrinkle

keep out or reach of the children

Take appropriate amount and evenly apply it to the skin

1. Open the lid of the vial and remove aluminum thread carefully
2. Insert vial in serter in the vial.
3. Drop the fomula on wrinkles around eyes and mouth and the entire face. Then use your fingertips to help formula to be absorbed into the skin.

If you experience following symptoms after using the cosmetics, you should immediately stop using the cosmetics. If you continue to use them, the symptoms may worsen. Consult with your dermatologist.
A) When you experience red spots, swelling, itchiness and irritation while applying the product.
B) When the applied area experience symptoms while it is exposed to direct sunlight
2. Do not use the cosmetics on the areas where you have wounds, eczema or dermatitis
3. Precautions when storing and using the product
A) Keep the lid closed after using the product
B) Store it out of reach of children
C) Do not store it in places with high or low temperature or where it is exposed to the sunlight directly.

for external use only

RUBY REBORN TRANSFUSION DNA KIT

glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70514-0004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	7 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ADENOSINE (UNII: K72T3FS567)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:70514-0004-1	22 mL in 1 AMPULE; Type 0: Not a Combination Product	12/05/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part347	11/29/2017	

Labeler - Karatica Co., Ltd (689605545)

Registrant - Karatica Co., Ltd (689605545)

Establishment

Name	Address	ID/FEI	Business Operations
Karatica Co., Ltd		689605545	manufacture(70514-0004)

Revised: 12/2017 Karatica Co., Ltd