PERSONAL LUBRICANT- sodium hyaluronate gel Jiangxi Renhetang pharmaceutical chain Co., Ltd.

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.

PERSONAL LUBRICANT

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Active Ingredient(s)

SODIUM HYALURONATE 0.01g/100mL

Purpose

BACTERIOSTATIC

Use

Uses for adult sexual wellness, which can produce lubrication to facilisexual activity.

Warnings

This product cannot be used as spermicide or contraceptives.

Please keep this product out of the reach of children.

Discontinue use should any allergy occur.

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

• Squeeze out the product and apply desired amount onto the intimate area. Reapply as desired. It can be used for vaginal sex, oral sex, anal sex, etc.

Other information

• Keep in sealed container, and place in a cool dry place.

Inactive ingredients

WATER, GLYCEROL, PROPYLENE GLYCOL, CELLULOSE, SODIUM HYALURONATE, ETHYL 2-HYDROXYBENZOATE, ALOE SERUM.

Package Label - Principal Display Panel



Expiration Date:

管径35mm





PERSONAL LUBRICA	NT				
sodium hyaluronate gel					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:81615-001	
Route of Administration	TOPICAL				
Active Ingredient/Active	Moiety				
Ingredient Name Basis o Strengt				Stra	ength
			HYALURONATE SODIUM	0.01 g in 100	mL
Inactive Ingredients					
	Ingredient Name			Streng	yth
ALOE VERA LEAF (UNII: ZY81Z83	H0X)				
POWDERED CELLULOSE (UNII: S	MD1X3XO9M)				
GLYCERIN (UNII: PDC6A3C0OX)					
PROPYLENE GLYCOL (UNII: 6DC9	Q167V3)				
WATER (UNII: 059QF0KO0R)					
ETHYL SALICYLATE (UNII: 555U6	ΓZ2MV)				

Packaging								
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:81615-001- 01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2021					
2	02	80 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2021					
3	NDC:81615-001- 03	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2021					
Μ	larketing	Information						
Μ	larketing Marketing Category	Information Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				

Labeler - Jiangxi Renhetang pharmaceutical chain Co., Ltd. (410551226)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Kangmei medical and Health Products Co., Ltd		528118770	manufacture(81615-001)

Revised: 4/2021

Jiangxi Renhetang pharmaceutical chain Co., Ltd.