QYO QYO TANGERINE BRIGHT MOIST PEELING PACK- kaolin paste
LaLa 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.

Qyo Qyo Tangerine Bright + Moist
Peeling Pack

ACTIVE INGREDIENTS

KAOLIN 5%

PURPOSE

SKIN PROTECTANT

USES

- Helps prevent and temporarily protects chapped or cracked skin.

WARNINGS

For external use only.

When using this product • do not get into eyes.

Stop us and ask doctor if • condition worsens • symptoms last more than 7 days or clear up and occurs again within a few days.

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

Directions

• Apply as needed. • After cleansing your face, apply two spatula of peeling pack and spread evenly on your face. • Wait 10 to 15 minutes. • Massage and deep cleanse your skin before rinsing with luke warm water.

Inactive Ingredients

Water, Sodium lauryl sulfate, Stearic Acid, Glycerin, Volcanic Ash, Buylene Glycol, Potassium Hydroxide, Bentonite, Betaine

Other Information

Protect this product from excessive heat and direct sun.

Questions or Comments?

Call toll free **1-800-311-6680**

PRINCIPAL DISPLAY PANEL

Qyo Qyo
Tangerine
Bright + Moist
Peeling Pack
100mL

QYO QYO TANGERINE BRIGHT MOIST PEELING PACK

kaolin paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71733-108

Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
KAOLIN (UNII: 24H4NWX5CO) (KAOLIN - UNII:24H4NWX5CO)	KAOLIN	5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
STEARIC ACID (UNII: 4ELV7Z65AP)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
TANGERINE PEEL (UNII: JU3D414057)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BENTONITE (UNII: A3N5ZCN45C)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71733-108-01	1 in 1 BOX	01/01/2018	
1		100 mL in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	01/01/2018	

Labeler - LaLa Co., Ltd. (694617781)

Establishment

Name	Address	ID/FEI	Business Operations
LaLa Co., Ltd.		694617781	manufacture(71733-108)

Revised: 1/2019 LaLa Co., Ltd.