

**V34 TOOTHPASTE. TABLETS.- v34 toothpaste tablets tablet  
LLRN PERSONAL CARE(SHENZHEN)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.*

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**Active Ingredient**

Sodium Bicarbonate 7%

Phthalimidoperoxyacaproic Acid 6%

Hydrated Silica 5%

**Purpose**

Whitening, Fresh Breath, Clean mouth

**Use**

Place tablets in your mouth.

Chew until it foams.

Brush your teeth as usual. Use twice daily.

**Warnings**

Products is not intended for use by children.

Do not swallow.

**Do not use**

Pregnant woman and children under 12 years old.

**When Using**

Do not swallow foams.

**Stop Use**

Stop use if you feel unwell or nausea.

**Ask Doctor**

Call a POISON CENTER or doctor/physician if you feel unwell for an extended period of time afer use.

# Keep Out Of Reach Of Children

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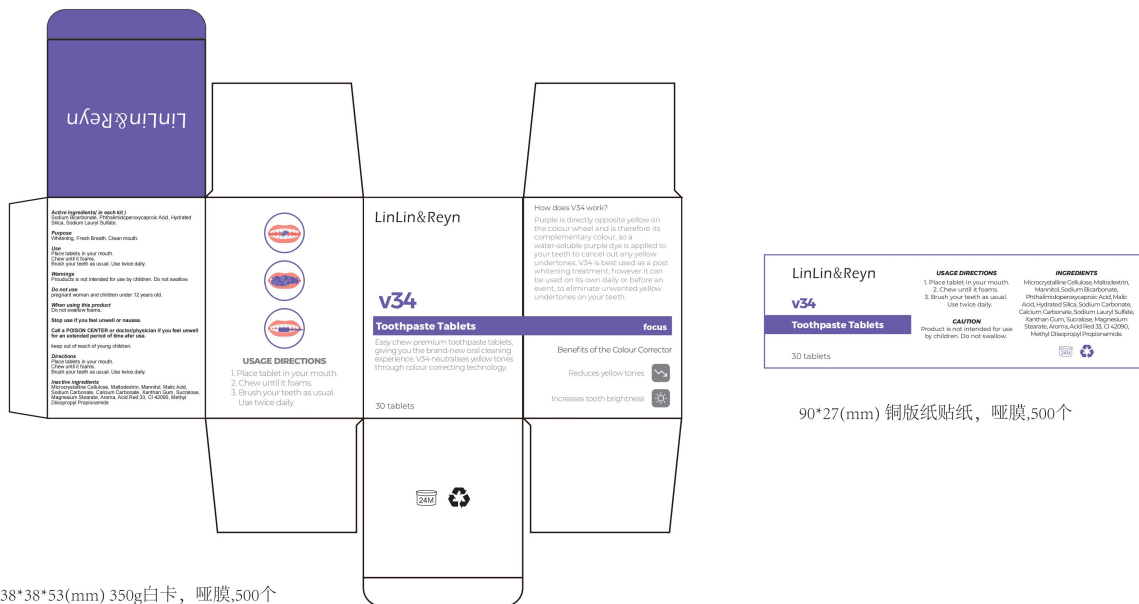
## Directions

Place tablets in your mouth.  
Chew until it foams.  
Brush your teeth as usual. Use twice daily.

## Inactive ingredients

Microcrystalline Cellulose, Maltodextrin, Mannitol, Malic Acid, Sodium Carbonate, Calcium Carbonate, Xanthan Gum, Sucralose, Magnesium Stearate, Aroma, Acid Red 33, CI 42090, Methyl Diisopropyl Propionamide

## PRINCIPAL DISPLAY PANEL



38\*38\*53(mm) 350g白卡, 哑膜, 500个

## V34 TOOTH PASTE. TABLETS.

v34 toothpaste tablets tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83812-005
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>PHTHALIMIDOPEROXYCAPROIC ACID</b> (UNII: 5OEJ6FAL6C) (PHTHALIMIDOPEROXYCAPROIC ACID - UNII:5OEJ6FAL6C)	PHTHALIMIDOPEROXYCAPROIC ACID	6 g in 100
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4MONH37)	SODIUM BICARBONATE	7 g in 100
<b>HYDRATED SILICA</b> (UNII: Y6O7T4G8P9) (HYDRATED SILICA - UNII:Y6O7T4G8P9)	HYDRATED SILICA	5 g in 100

## Inactive Ingredients

Ingredient Name	Strength
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>METHYL DIISOPROPYL PROPIONAMIDE</b> (UNII: 6QOP5A9489)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>AROMADENDRIN</b> (UNII: 7YA4640575)	
<b>SODIUM CARBONATE</b> (UNII: 45P3261C7T)	
<b>MALIC ACID</b> (UNII: 817L1N4CKP)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	5mm
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83812-005-01	30 in 1 BOTTLE; Type 0: Not a Combination Product	11/24/2023	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/24/2023	

**Labeler** - LLRN PERSONAL CARE(SHENZHEN)CO., LTD (419890311)

## **Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
LLRN PERSONAL CARE(SHENZHEN)CO., LTD		419890311	label(83812-005) , manufacture(83812-005)

Revised: 11/2023

LLRN PERSONAL CARE(SHENZHEN)CO., LTD