

**NICOTINE GUM- nicotine polacrilex gum, chewing**  
**Guangzhou Shuimu Jiahe Trading Co., Ltd.**

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reduces withdrawal symptoms, including nicotine craving, associated with quitting smoking

Microcrystalline cellulose, mannitol, propylene glycol sodium chloride, sodium bicarbonate, acesulfame, edible essence.

Nicotine (8 mg per pouch)

Ask a doctor before use if you have

a sodium-restricted diet

heart disease, recent heart attack, or irregular heartbeat. Nicotine can increase your heart rate.

high blood pressure not controlled with medication. Nicotine can increase blood pressure.

Ask a doctor or pharmacist before use if you are

using a non-nicotine stop smoking drug

taking prescription medicine for depression or asthma. Your prescription dose may need to be adjusted.

Stop use and ask a doctor if

mouth, teeth or jaw problems occur

irregular heartbeat or palpitations occur

you get symptoms of nicotine overdose such as nausea, vomiting, dizziness, diarrhea, weakness and rapid heartbeat

you have symptoms of an allergic reaction (such as difficulty breathing or rash).

Keep out of reach of children

Stop smoking aid

If you are pregnant or breast-feeding, only use this medicine on the advice of your health care provider. Smoking can seriously harm your child. Try to stop smoking without using any nicotine replacement medicine. This medicine is believed to be safer than smoking. However, the risks to your child from this medicine are not fully known.

if you are under 18 years of age, ask a doctor before use

before using this product, read the enclosed User's Guide for complete directions and other important information

begin using the gum on your quit day

if you smoke your first cigarette more than 30 minutes after waking up, use Nicotine Polacrilex Gum, 4 mg

your first cigarette within 30 minutes of waking up, use Nicotine Polacrilex Gum, 4 mg



## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:87494-002
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>NICOTINE</b> (UNII: 6M3C89ZY6R) (NICOTINE - UNII:6M3C89ZY6R)	NICOTINE	4 mg in 1000 mg

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>SODIUM CARBONATE</b> (UNII: 45P3261C7T)	
<b>CARNAUBA WAX</b> (UNII: R12CBM0EIZ)	
<b>ACESULFAME POTASSIUM</b> (UNII: 23OV73Q5G9)	
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>MINT</b> (UNII: FV98Z8GITP)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>XYLITOL</b> (UNII: VCQ006KQ1E)	
<b>SORBITOL</b> (UNII: 506T60A25R)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	score with uneven pieces
<b>Shape</b>	SQUARE	<b>Size</b>	10mm
<b>Flavor</b>	MINT	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:87494-002-01	48 mg in 1 BOX, UNIT-DOSE; Type 0: Not a Combination Product	04/03/2026	

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012	04/03/2026	

**Labeler** - Guangzhou Shuimu Jiahe Trading Co., Ltd. (700938261)

**Establishment**

Name	Address	ID/FEI	Business Operations
Guangzhou Shuimu Jiahe Trading Co., Ltd.		700938261	manufacture(87494-002)

Revised: 4/2026

Guangzhou Shuimu Jiahe Trading Co., Ltd.