NICORETTE- nicotine polacrilex lozenge Haleon US Holdings LLC

Drug Facts

Active ingredient (in each lozenge) (2 mg)

Nicotine polacrilex, 2 mg

Active ingredient (in each lozenge) (4 mg)

Nicotine polacrilex, 4 mg

Purpose

Stop smoking aid

Use

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

Warnings

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.

Do not use (Mint)

• if you are allergic to soya

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 your blood pressure.
- stomach ulcer or diabetes
- history of seizures

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 problems occur
- persistent indigestion or severe sore throat occurs
- 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 and pets.

Nicotine lozenges may have enough nicotine to make children and pets sick. If you need to remove the lozenge, wrap it in paper and throw away in the trash. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions (2 mg)

- if you are under 18 years of age, ask a doctor before use. No studies have been done to show if this product will work for you.
- before using this product, read the enclosed User's Guide for complete directions and other important information
- begin using the lozenge on your quit day
- if you smoke your first cigarette within 30 minutes of waking up, use 4 mg nicotine lozenge
- if you smoke your first cigarette more than 30 minutes after waking up,use 2 mg nicotine lozenge according to the following 12 week schedule:

Weeks 1 to 6	Weeks 7 to 9	Weeks 10 to 12
1 lozenge every 1 to 2 hours	1 lozenge every 2 to 4 hours	1 lozenge every 4 to 8 hours

- nicotine lozenge is a medicine and must be used a certain way to get the best results
- place the lozenge in your mouth and allow the lozenge to slowly dissolve (about 20 30 minutes). Minimize swallowing. **Do not chew or swallow lozenge.**
- you may feel a warm or tingling sensation
- occasionally move the lozenge from one side of your mouth to the other until completely dissolved (about 20 - 30 minutes)
- do not eat or drink 15 minutes before using or while the lozenge is in your mouth
- to improve your chances of quitting, use at least 9 lozenges per day for the first 6 weeks
- do not use more than one lozenge at a time or continuously use one lozenge after another since this may cause you hiccups, heartburn, nausea or other side effects
- do not use more than 5 lozenges in 6 hours. Do not use more than 20 lozenges per day.
- it is important to complete treatment. If you feel you need to use the lozenge for a longer period to keep from smoking, talk to your health care provider.

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Other information

- each lozenge contains:sodium, 18 mg
- Phenylketonurics: Contains Phenylalanine 3.4 mg per lozenge
- store at 20 25°C (68 77°F)
- keep POPPAC tightly closed and protect from light

Inactive ingredients

acacia, aspartame, calcium polycarbophil, corn syrup solids, flavors, lactose, magnesium stearate, maltodextrin, mannitol, potassium bicarbonate, sodium alginate, sodium carbonate, soy protein, triethyl citrate, xanthan gum

Questions or comments?

call toll-free **1-888-569-1743**

Principal Display Panel

NDC 0135-0510-07

Nicorette ®

nicotine polacrilex lozenge, 2 mg stop smoking aid

Lozenge

Includes User's Guide

2mg

FOR THOSE WHO SMOKE THEIR FIRST CIGARETTE MORE THAN 30 MINUTES AFTER WAKING UP.If you smoke your first cigarette <u>WITHIN</u>30 MINUTES of waking up, use Nicorette 4 mg Lozenge

Mint

144 Lozenges,2 mg Each

(6 POPPAC ® Containers of 24)

- not for sale to those under 18 years of age
- proof of age required
- not for sale in vending machines or from any source where proof of age cannot be verified

TAMPER EVIDENT FEATURE: Do not use if clear neckband printed "SEALED FOR SAFETY" is missing or broken.

Retain outer carton for full product uses, directions and warnings.

TO INCREASE YOUR SUCCESS IN QUITTING:

- 1. You must be motivated to quit.
- 2. **Use Enough -**Use **at least 9 Nicorette** [®]Lozenges per day during the first six weeks.
- 3. **Use Long Enough -**Use **Nicorette** ®Lozenges for the full 12 weeks.
- 4. **Use With a Support Program**as directed in the enclosed User's Guide.

Nicorette ® POPPAC ®

To open vial, push in child resistant band on the POPPAC [®] container with thumb.

Flip up the top of the POPPAC $^{\$}$ and remove lozenge. A small amount of powder on opening of the POPPAC $^{\$}$ is normal.

For more information and for a FREE individualized stop smoking program, please visit www.Nicorette.comor see inside for more details.

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Distributed By:

GSKConsumer Healthcare

Warren, NJ 07059

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Principal Display Panel
NDC 0135-0511-07
Nicorette ®

nicotine polacrilex lozenge, 4 mg stop smoking aid

Lozenge

Includes User's Guide

4 mg

FOR THOSE WHO SMOKE THEIR FIRST CIGARETTE WITHIN 30 MINUTES OF WAKING UP.

If you smoke your first cigarette **MORE THAN30 MINUTES** after waking up, use Nicorette® 2 mg Lozenge

Mint

144 Lozenges,4 mg Each

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For more information and for a FREE individualized stop smoking program, please visit <u>www.Nicorette.com</u>or see inside for more details.

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Distributed By:

GSKConsumer Healthcare

Warren, NJ 07059

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NICORETTE nicotine polacrilex lozenge			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-0510
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
NICOTINE (UNII: 6M3C89ZY6R) (NICOTINE - UNII:6M3C89ZY6R)	NICOTINE	2 mg

Inactive Ingredients	
Ingredient Name	Strength
POLACRILIN (UNII: RCZ785HI7S)	
ACACIA (UNII: 5C5403N26O)	
ASPARTAME (UNII: Z0H242BBR1)	
CALCIUM POLYCARBOPHIL (UNII: 8F049NKY49)	
CORN SYRUP (UNII: 9G5L16BK6N)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MANNITOL (UNII: 30WL53L36A)	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
SOY PROTEIN (UNII: R44IWB3RN5)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	white (white to off-white)	Score	no score
Shape	ROUND	Size	16mm
Flavor	MINT	Imprint Code	NL2S
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0510- 01	3 in 1 CARTON	10/19/2010	
1		24 in 1 CONTAINER; Type 0: Not a Combination Product		
2	NDC:0135-0510- 06	1 in 1 CARTON	09/01/2015	
2		24 in 1 CONTAINER; Type 0: Not a Combination Product		
3	NDC:0135-0510- 07	6 in 1 CARTON	09/01/2015	
3		24 in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021330	10/19/2010	

NICORETTE

nicotine polacrilex lozenge

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0135-0511

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NICOTINE (LINII) 6M3C807Y6P) (NICOTINE LINII) 6M3C807Y6P)	NICOTINE	1 ma

Inactive Ingredients

Ingredient Name	Strength
POLACRILIN (UNII: RCZ785HI7S)	
ACACIA (UNII: 5C5403N26O)	
ASPARTAME (UNII: Z0H242BBR1)	
CALCIUM POLYCARBOPHIL (UNII: 8F049NKY49)	
CORN SYRUP (UNII: 9G5L16BK6N)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MANNITOL (UNII: 3OWL53L36A)	
POTASSIUM BICARBONATE (UNII: HM5Z15LEBN)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
SOY PROTEIN (UNII: R44IWB3RN5)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	

Product Characteristics

XANTHAN GUM (UNII: TTV12P4NEE)

Color	white (white to off-white)	Score	no score
Shape	ROUND	Size	16mm
Flavor	MINT	Imprint Code	NL4S
Contains			

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-					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
п					

1	01 NDC:0132-0211-	3 in 1 CARTON	10/19/2010
1		24 in 1 CONTAINER; Type 0: Not a Combination Product	
2	NDC:0135-0511- 06	1 in 1 CARTON	09/01/2015
2		24 in 1 CONTAINER; Type 0: Not a Combination Product	
3	NDC:0135-0511- 07	6 in 1 CARTON	09/01/2015
3		24 in 1 CONTAINER; Type 0: Not a Combination Product	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
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Labeler - Haleon US Holdings LLC (079944263)

Revised: 3/2024 Haleon US Holdings LLC