

**BISMUTH SUBSALICYLATE- sunmark stomach relief tablet, chewable
NuCare Pharmaceuticals, Inc.**

Sunmark Stomach relief tablets Regular Strength

Active ingredient (in each tablet)

Bismuth subsalicylate 262 mg

(total salicylate 102 mg per tablet)

Purpose

Upset stomach reliever and anti-diarrheal

Uses

relieves:

- diarrhea
- heartburn
- indigestion
- nausea
- upset stomach associated with these symptoms

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Contains salicylate. Do not take if you are

- allergic to salicylates (including aspirin)
- taking other salicylate products

Do not use if you have

- bloody or black stools
- an ulcer
- a bleeding problem

Ask a doctor before use if you have

- fever
- mucus in the stool

Ask a doctor or pharmacist before use if you are

- taking any drug for

- anticoagulation (thinning of the blood)
- diabetes
- gout
- arthritis

When using this product

a temporary, but harmless darkening of the stool and/or tongue may occur.

Stop use and ask a doctor if

- symptoms get worse
- ringing in the ears or loss of hearing occurs
- diarrhea lasts more than 2 days

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center immediately.

Directions

- chew or dissolve in mouth
- adults and children 12 years and over: 2 tablets every 1/2 to 1 hour as needed
- do not take more than 8 doses (16 tablets) in 24 hours
- children under 12 years: ask a doctor
- drink plenty of fluids to help prevent dehydration which may accompany diarrhea.

Other information

each tablet contains:

- Sodium less than 1 mg
- salicylate 102 mg
- very low sodium
- avoid excessive heat (over 104°F or 40°C)
- TAMPER EVIDENT: Do not use if individual compartments are torn or missing.

Inactive ingredients

calcium carbonate, D&C red 27 aluminum lake, flavor, magnesium stearate, mannitol, pregelatinized starch, saccharin sodium.

Principal Display Panel

Distributed by:
 McKesson San Francisco, CA
 94104
 Packaged By:
 NuCare Pharmaceuticals, Inc.
 Orange, CA 92667
 Patient Instructions:
 Chew _____ times a day,
 every _____ hours
 68071452303*30*000000*000000
 Rev 01/01/19

NDC: 68071-4523-3
Stomach Relief Tablets
#30 Chewtabs

Bismuth Subsalicylate 262mg
 (total Salicylate 102mg per tablet)
 See manufacturer's label
 for full list of ingredients.

Stomach Relief Tablets
 Lot: 000000 NDC: 68071-4523-03
 MFR NDC: 49348-953-44 Exp.: 00-00
 Serial# 00000000002

Stomach Relief Tablets
 Lot: 000000 NDC: 68071-4523-03
 MFR NDC: 49348-953-44 Exp.: 00-00
 Serial# 00000000002



GTIN 00368071452334
 Serial# 00000000002
 Exp. Date 00-00
 LOT#: 000000

Call your doctor for medical advice about side
 effects. You may report side effects to FDA at
 1-800-FDA-1088.

Product #: R0657030

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 59-86°F.

BISMUTH SUBSALICYLATE

sunmark stomach relief tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4523(NDC:49348-953)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	262 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
STARCH, CORN (UNII: O8232NY3SJ)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

Product Characteristics

Color	pink	Score	no score
Shape	ROUND	Size	17mm
Flavor		Imprint Code	GDC122
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4523-3	30 in 1 BOX; Type 0: Not a Combination Product	08/10/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	03/01/2011	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-4523)

Revised: 6/2024

NuCare Pharmaceuticals,Inc.