

SENOKOT- standardized senna concentrate tablet, film coated
Atlantis Consumer Healthcare, Inc.

SenokotXTRA
(standardized senna concentrate)

Drug Facts

Active ingredient (in each tablet) Purpose

Sennosides 17.2 mg

Purpose

Laxative

Uses

- relieves occasional constipation (irregularity)
- generally causes bowel movement in 6-12 hours

Warnings

Do not use

- laxative products for longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel movements that continues over a period of 2 weeks

Stop use and ask a doctor if you have rectal bleeding or fail to have a bowel movement after use of a laxative. These may indicate a serious condition.

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 right away.

Directions

- take preferably at bedtime or as directed by a doctor

age	starting dosage	maximum dosage
adults and children 12 years of age and over	1 tablet once a day	2 tablets twice a day
children 6 to under 12 years	½ tablet once a day	1 tablet twice a day
children under 6	ask a doctor	ask a doctor

Other information

- each tablet contains: **calcium 20 mg**

- store at 25°C (77°F); excursions permitted between 15°-30°C(59°-86°F)

Inactive ingredients croscarmellosesodium, dicalcium phosphate, hypromellose, lactose, magnesium stearate, microcrystalline cellulose, mineral oil, stearic acid, talc, tartaric acid

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Dist. by: **Avrio Health L.P., Stamford, CT 06901-3431**

Senokot® Extra Strength
36 Tablets Carton



Drug Facts (continued)

Directions

take preferably at bedtime or as directed by a doctor

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adults and children 12 years and over	1 tablet once a day	2 tablets twice a day
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Senokot and the Comfort Promise Seal are registered trademarks of Atlantis Consumer Healthcare Inc.

A1023

The Senokot® Laxative Comfort Promise®

For your comfort, the active ingredient in Senokot laxative tablets is always purified senna, manufactured to high quality standards.



Dist. by: Atlantis Consumer Healthcare Inc. Bridgewater, NJ 08807 USA
 Questions or comments? 1-833-288-2684
 www.senokot.com
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Senokot®
Standardized Senna Concentrate

Extra Strength

Senokot®
Standardized Senna Concentrate, 17.2mg

Extra Strength

Natural vegetable laxative ingredient

Extra strength, just as gentle

Dependable overnight constipation relief

36 Tablets



Drug Facts

Active ingredient Purpose (in each tablet)
 Sennosides 17.2 mg.....Laxative

Uses

relieves occasional constipation (irregularity) generally produces a bowel movement in 6 to 12 hours

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Do not use if seal under cap is missing or damaged.

R52519

Senokot® Extra Strength
12Tablets Carton

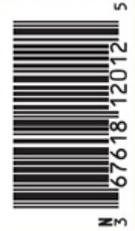
Senokot[®]
Standardized Senna Concentrate

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Standardized Senna Concentrate, 17.2mg

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Extra Strength

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Inactive ingredients croscarmellose sodium, dicalcium phosphate, hypromellose, lactose, magnesium stearate, microcrystalline cellulose, mineral oil, stearic acid, talc, tartaric acid

Tamper Evident: Do not use if sealed blister unit is broken or damaged. Retain this carton for important information.

R52517

Senokot[®] Extra Strength 36 Tablets Label



SENOKOT

standardized senna concentrate tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67618-120
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	17.2 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TARTARIC ACID (UNII: W4888I119H)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
TALC (UNII: 7SEV7J4R1U)	
STEARIC ACID (UNII: 4ELV7Z 65AP)	

Product Characteristics

Color	BROWN (Light Brown)	Score	no score
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Shape	ROUND	Size	9mm
Flavor		Imprint Code	X
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67618-120-12	1 in 1 CARTON	09/01/1988	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:67618-120-36	3 in 1 CELLO PACK	09/01/1988	01/01/2023
2		12 in 1 BLISTER PACK		
2		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:67618-120-06	36 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/1988	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	09/01/1988	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Registrant - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 12/2023

Atlantis Consumer Healthcare, Inc.