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

Senokot
(standardized senna concentrate)

Drug Facts

Active ingredient (in each tablet)

Sennosides 8.6mg

Purpose

Laxative

Uses

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

Warnings

Do not use

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

Aska doctor before use if you have

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

Stop use and ask a doctorif you have rectal bleeding or fail to have a bowel movementafter 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 ControlCenter 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	2 tablets oncea day	4 tablets twicea day
children 6 to under12 years	1 tablet once aday	2 tablets twicea day
children 2 to under6 years	1/2 tablet oncea day	1 tablet twicea day
children under2 years	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

croscarmellose sodium, dicalciumphosphate, hypromellose, magnesium stearate, maltodextrin, microcrystallinecellulose, mineral oil, polyethylene glycol, talc

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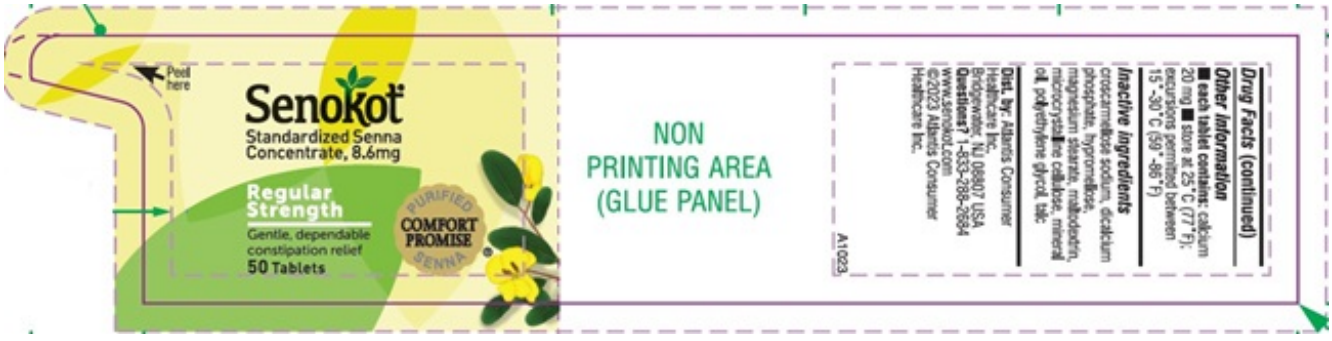
Avrio Health L.P.

305616-0B

50 Tablets
NDC 67618-300-50



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

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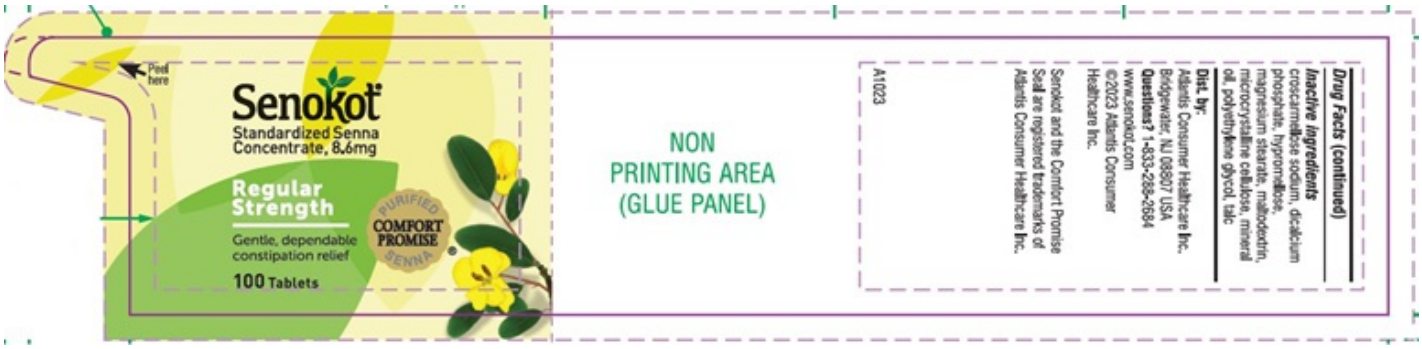
adults and children 12 years of age and over	2 tablets once a day	4 tablets twice a day
children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
children 2 to under 6 years	½ tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

100 Tablets
NDC 67618-300-10



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SENOKOT

standardized senna concentrate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67618-300

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	BROWN	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	S
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67618-300-20	2 in 1 CARTON	06/01/1956	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:67618-300-50	1 in 1 CARTON	06/01/1956	
2		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:67618-300-10	1 in 1 CARTON	06/01/1956	
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	06/01/1956	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Registrant - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 12/2025 Atlantis Consumer Healthcare, Inc.