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

SenokotXTRA
(standardizedsenna 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 directedby a doctor

Ask a doctor beforeuse if you have

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

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

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

Keep out of reachof children. In case of overdose, get medical help or contacta Poison Control Center right away.

Directions

- take preferably at bedtime or as directed by a doctor

age	startingdosage	maximumdosage
adults and children 12 yearsof age and over	1 tablet once a day	2 tablets twice aday
children 6 to under 12 years	½ tablet once a day	1 tablet twice aday
children under 6	ask a doctor	ask a doctor

Other information

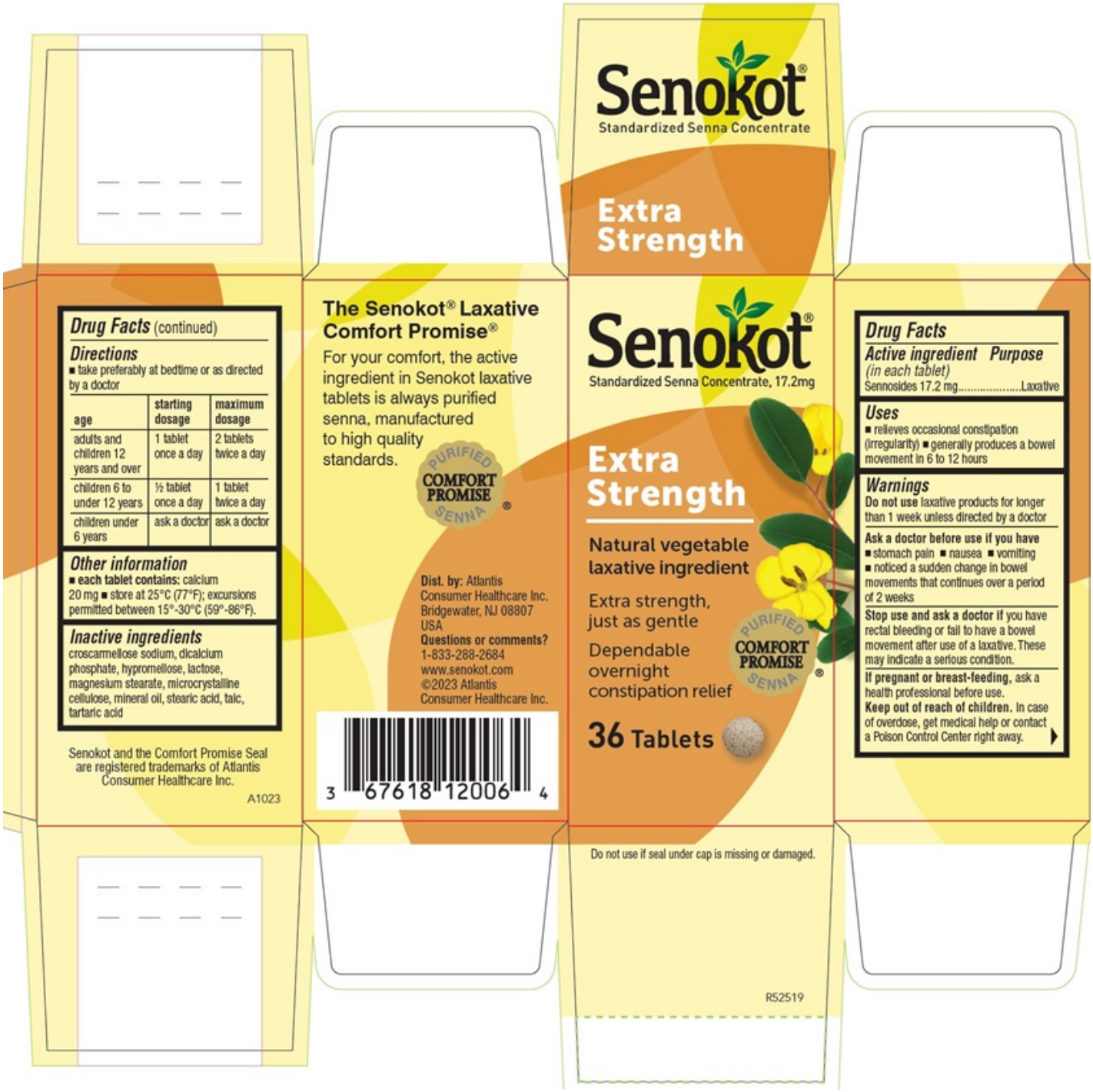
- each tablet contains: **calcium 20 mg**

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

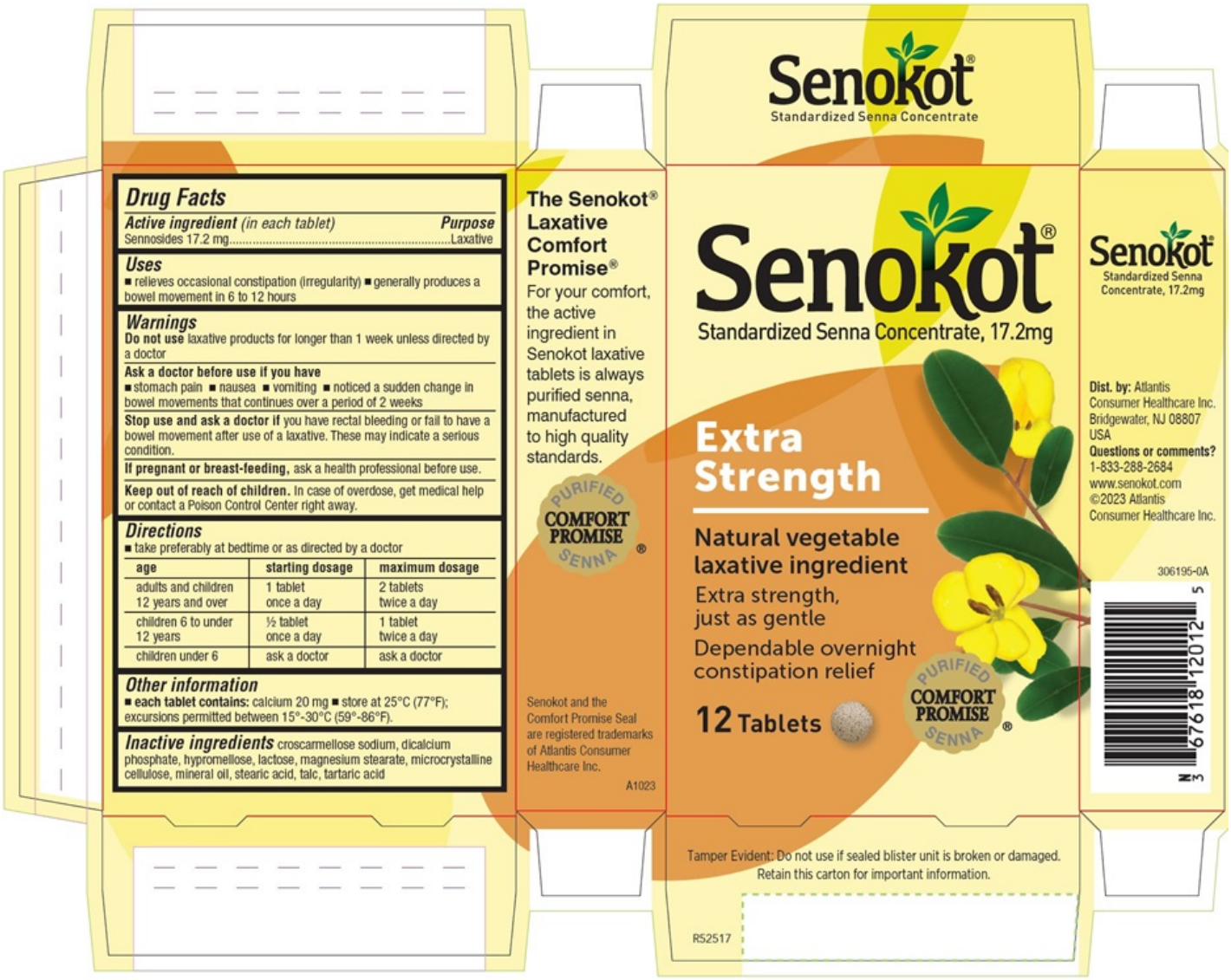
Inactive ingredients croscarmellosesodium, dicalcium phosphate, hypromellose, lactose, magnesium stearate,microcrystalline cellulose, mineral oil, stearic acid, talc, tartaricacid

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

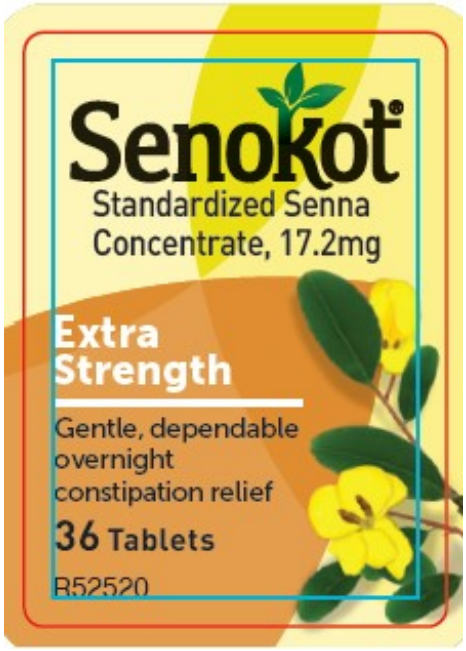
Senokot® Extra Strength
36 Tablets Carton



Senokot® Extra Strength
12Tablets Carton



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**
CROSCARMELLOSE 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: 4ELV7Z65AP)	

Product Characteristics			
Color	BROWN (Light Brown)	**Score**	no score

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.