DOCUSATE SODIUM AND SENNOSIDES- docusate sodium and sennosides capsule, liquid filled
SOFTGEL HEALTHCARE PRIVATE LIMITED

DOCUSATE SODIUM 50 MG & SENNOSIDE 8.6 MG CAPSULES (Red)

Active ingredient (in each softgel)

Docusate Sodium 50 mg

Sennosides 8.6 mg

Purpose

Stool softener

Laxative

Uses

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

Warnings
Do not use

- if you are now taking mineral oil, unless directed by a doctor
- 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 be signs of a serious condition.

If pregnant or breast-feeding

ask a health professional before use.

Keep out of reach of children

In case of accidental overdose, seek professional assistance or contact a Poison Control Center right away.

Directions

- take preferably at bedtime or as directed by a doctor.

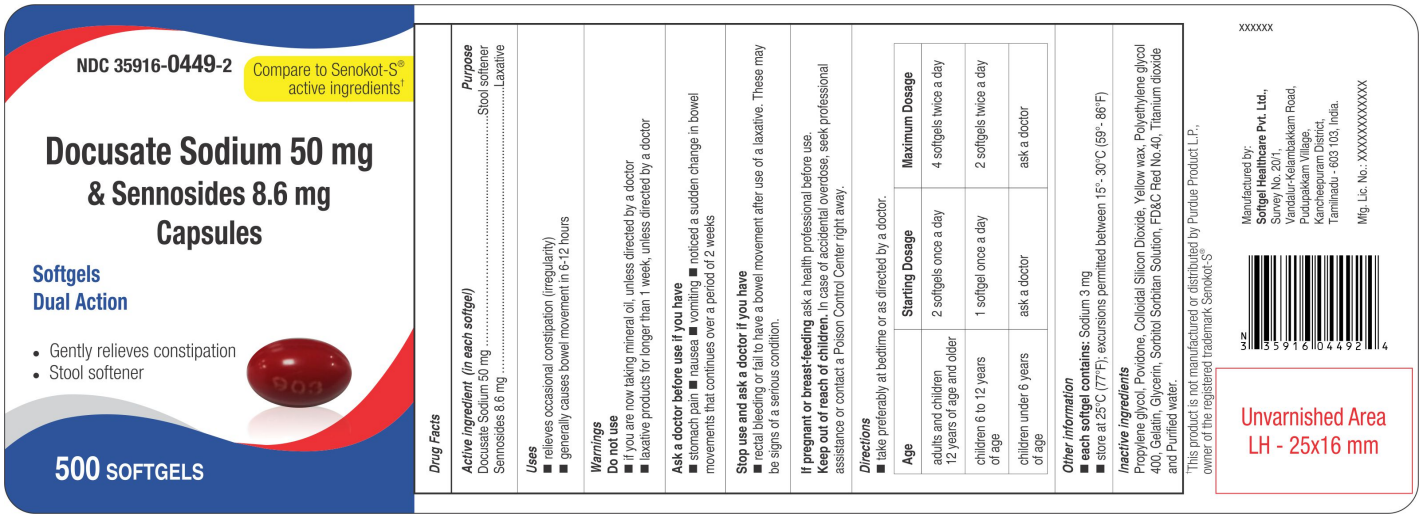
Age	Starting Dosage	Maximum Dosage
adults and children 12 years of age and older	2 softgels once a day	4 softgels twice a day
children 6 to 12 years of age	1 softgel once a day	2 softgels twice a day
children under 6 years of age	ask a doctor	ask a doctor

Other information

- **each softgel contains:** Sodium 3 mg
- store at 25°C (77°F); excursions permitted between 15°- 30°C (59°- 86°F)

Inactive ingredients

Propylene glycol, Povidone, Colloidal Silicon Dioxide, Yellow wax, Polyethylene glycol 400, Gelatin, Glycerin, Sorbitol Sorbitan Solution, FD&C Red No.40, Titanium dioxide and Purified water.



DOCUSATE SODIUM AND SENNOSIDES

docusate sodium and sennosides capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:35916-0449
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg

Inactive Ingredients

Ingredient Name	Strength
POVIDONE K30 (UNII: U725QWY32X)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SORBITOL (UNII: 506T60A25R)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
1,4-SORBITAN (UNII: AV0YTZ4E6J)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	red ((Opaque))	**Score**	score with uneven pieces
Shape	OVAL	**Size**	5mm
Flavor		**Imprint Code**	903
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:35916-0449-2	500 in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	04/22/2024	

Labeler - SOFTGEL HEALTHCARE PRIVATE LIMITED (675584180)

Establishment

Name	Address	ID/FEI	Business Operations
SOFTGEL HEALTHCARE PRIVATE LIMITED		675584180	manufacture(35916-0449)