SENNA WITH DOCUSATE SODIUM- docusate sodium and sennosides tablet
HHH Pharmaceuticals

Drug Facts

Active ingredients (in each tablet) ***Purpose***

Docusate sodium 50 mg.....Stool softner

Sennosides 8.6 mg.....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 habits 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	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

Other information

- each tablet contains: calcium 21 mg, sodium 3 mg. VERY LOW SODIUM
- store at 25ºC (77ºF); excursions permitted between 15°-30°C (59°-86°F)
- **DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

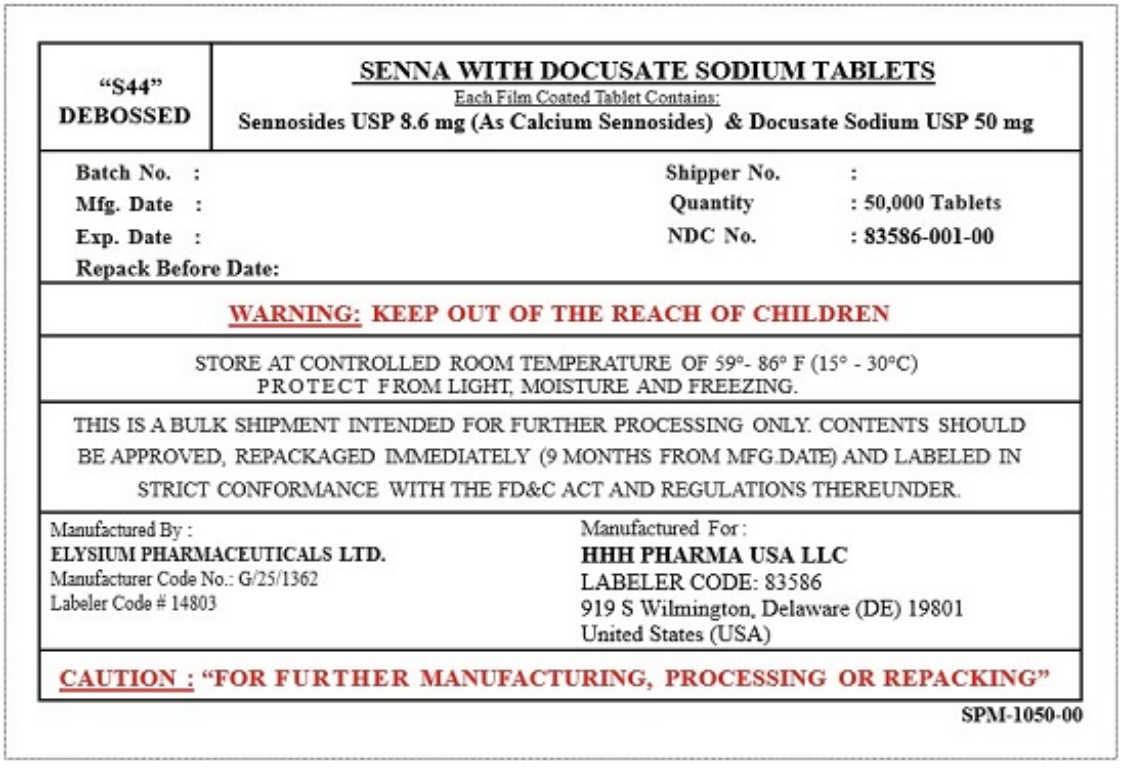
Inactive ingredients

croscarmellose sodium, colloidal silicon dioxide, dicalcium phosphate, polyethylene glycol, FD&C red 40 lake, hypromellose, magnesium stearate, microcrystalline cellulose, stearic acid powder, titanium dioxide.

Questions or comments?

call toll-free 1-302-380-3408

PRINCIPAL DISPLAY PANEL



"S44" DEBOSSED	**SENNA WITH DOCUSATE SODIUM TABLETS** Each Film Coated Tablet Contains: **Sennosides USP 8.6 mg (As Calcium Sennosides) & Docusate Sodium USP 50 mg**

Batch No. : **Shipper No. :**
Mfg. Date : **Quantity : 50,000 Tablets**
Exp. Date : **NDC No. : 83586-001-00**
Repack Before Date:

WARNING: KEEP OUT OF THE REACH OF CHILDREN

STORE AT CONTROLLED ROOM TEMPERATURE OF 59°- 86° F (15° - 30°C)
PROTECT FROM LIGHT, MOISTURE AND FREEZING.

THIS IS A BULK SHIPMENT INTENDED FOR FURTHER PROCESSING ONLY. CONTENTS SHOULD BE APPROVED, REPACKAGED IMMEDIATELY (9 MONTHS FROM MFG.DATE) AND LABELED IN STRICT CONFORMANCE WITH THE FD&C ACT AND REGULATIONS THEREUNDER.

Manufactured By :
ELYSIUM PHARMACEUTICALS LTD.
Manufacturer Code No.: G/25/1362
Labeler Code # 14803

Manufactured For :
HHH PHARMA USA LLC
LABELER CODE: 83586
919 S Wilmington, Delaware (DE) 19801
United States (USA)

CAUTION : "FOR FURTHER MANUFACTURING, PROCESSING OR REPACKING"

SPM-1050-00

SENNA WITH DOCUSATE SODIUM

docusate sodium and sennosides tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83586-001

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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	red	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	S44
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83586-001-00	50000 in 1 DRUM; Type 0: Not a Combination Product	10/07/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/07/2025	

Labeler - HHH Pharmaceuticals (144848997)

Registrant - HHH Pharmaceuticals (144848997)

Establishment

Name	Address	ID/FEI	Business Operations
Elysium Pharmaceuticals Ltd		863182240	manufacture(83586-001)

Establishment

Name	Address	ID/FEI	Business Operations
HHH Pharmaceuticals		144848997	pack(83586-001)

Revised: 10/2025 HHH Pharmaceuticals