

DOCUSATE SODIUM AND SENNOSIDES- docusate sodium and
sennosides tablet
Morepen Laboratories Limited

Docusate Sodium 50 mg and Sennosides 8.6 mg Tablets

Active Ingredients Label

Drug Facts

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

Docusate sodium USP

50mg Stool softner

Sennosides 8.6 mg..... Laxative

Inactive Ingredients Section Label

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide.

*contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.

Purpose Section Label

Drug Facts

Active ingredient (in each tablet) Purpose

Docusate sodium USP

50mg Stool softner

Sennosides 8.6 mg..... Laxative

Indications & Usage Seciton Label

Uses

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

Dosage & Administration Section Label

Directions

■ take preferably at bedtime
or as directed by a doctor

age	Starting dosage	maximum dosage
adults and children 12 years 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	$\frac{1}{2}$ tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

Questions Section Label

Question or comments?

Call toll- free 1-844-317-6549 Mon-Fri 9AM-5PM
EST or contact the FDA at 1-800-FDA-1088 or
www.fda.gov/medwatch

Warnings Section Label

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 movement 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.



Keep out of reach of children.

In case of overdose, get
medical help or contact a
Poison Control Center
right away.



Storage and Handling Section Label

Other information

■ **each tablet contains:**

calcium 17.7 mg, sodium
4.4 mg VERY LOW SODIUM

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

30 Tablets Bottle Label (NDC 62993-0002-1)

NDC 62993-0002-1

Distributed by :
Morepen Bio Inc.
666, Plainsboro Road,
Suite 215 Plainsboro, NJ 08536

Dr. Morepen®

Docusate Sodium 50 mg & Sennosides 8.6 mg Tablets

Standardized Senna Concentrate, 8.6 mg
Docusate Sodium, 50 mg

*Compare to active ingredient in Senokot®-S

Dual Action

Gentle, dependable constipation relief

30 Tablets

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

No varnish Area for Lot No., Exp Date 15(L) x 24(H) mm

3 62993 00021 5 Peel here

Drug Facts

Active ingredient (in each tablet) Purpose

Docusate sodium USP 50mgStool softner
Sennosides 8.6 mg.... Laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces a 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 movement 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.

Drug Facts (continued)

Directions

■ take preferably at bedtime or as directed by a doctor

age	Starting dosage	maximum dosage
adults and children 12 years 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 17.7 mg, sodium 4.4 mg VERY LOW SODIUM
- store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide.

*contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.

Question or comments?

Call toll-free 1-844-317-6549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

* This product is not manufactured or distributed by Atlantis consumer healthcare Inc., owner of the registered trademark Senokot®-S

60 Tablets Bottle Label (NDC 62993-0002-2)

NDC 62993-0002-2



Docusate Sodium 50 mg & Sennosides 8.6 mg Tablets

Standardized Senna Concentrate, 8.6 mg

Docusate Sodium, 50 mg

*Compare to active ingredient in Senokot®-S

Dual Action

Gentle, dependable constipation relief

60 Tablets

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



N 3

6 2 9 9 3 1 0 0 2 2 2

No varnish Area for Lot No., Exp Date 15(L) x 24(H) mm

Distributed by :
Morepen Bio Inc.
666, Plainsboro Road,
Suite 215 Plainsboro, NJ 08536

Peel here

Drug Facts	
Active ingredient (in each tablet)	Purpose
Docusate sodium USP 50mgStool softner	
Sennosides 8.6 mg..... Laxative	
Uses	
- relieves occasional constipation (irregularity) - generally produces a 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 movement 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.	

Drug Facts (continued)			
Directions			
take preferably at bedtime or as directed by a doctor			
age	Starting dosage	maximum dosage	
adults and children 12 years 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 17.7 mg, sodium 4.4 mg VERY LOW SODIUM - store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).			
Inactive ingredients			
colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide.			
*contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.			
Question or comments?			
Call toll- Free 1-844-317-6549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch			
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100 Tablets Bottle Label (NDC 62993-0002-3)

NDC 62993-0002-3



Docusate Sodium 50 mg & Sennosides 8.6 mg Tablets

Standardized Senna Concentrate, 8.6 mg
Docusate Sodium, 50 mg

*Compare to active ingredient
in Senokot®-S

Dual Action

Gentle, dependable constipation relief

100 Tablets

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

No varnish Area
for Lot No., Exp Date
17(L) x 25(H) mm

Distributed by :
Morepen Bio Inc.
666, Plainsboro Road,
Suite 215 Plainsboro, NJ 08536



Peel here

Drug Facts	
Active ingredient (in each tablet)	Purpose
Docusate sodium USP 50mgStool softner Sennosides 8.6 mgLaxative	
Uses	
■ relieves occasional constipation (irregularity) ■ generally produces a 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 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.	

Drug Facts (continued)		
Directions		
■ take preferably at bedtime or as directed by a doctor		
age	Starting dosage	maximum dosage
adults and children 12 years 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 17.7 mg, sodium 4.4 mg VERY LOW SODIUM ■ store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).		
Inactive ingredients		
colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide. *contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.		
Question or comments?		
Call toll-free 1-844-317-6549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch		
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1000 Tablets Bottle Label (NDC 62993-0002-4)

NDC 62993-0002-4

Drug Facts**Active ingredient**

(in each tablet) **Purpose**
 Docusate sodium USP 50mg Stool softener
 Sennosides 8.6 mg Laxative

Uses ■ relieves occasional constipation (irregularity)
 ■ generally produces a 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.



Docusate Sodium 50 mg & Sennosides 8.6 mg Tablets

Standardized Senna Concentrate, 8.6 mg
 Docusate Sodium, 50 mg

*Compare to active ingredient in Senokot®-S

Dual Action

Gentle, dependable constipation relief

1000 Tablets

Drug Facts (continued)

Directions ■ take preferably at bedtime or as directed by a doctor

age	Starting dosage	maximum dosage
adults and children 12 years 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 17.7 mg, sodium 4.4 mg VERY LOW SODIUM ■ store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).

Inactive ingredients colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide. *contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.

Question or comments? Call toll-free 1-844-317-8549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

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

No varnish Area for Lot No., Exp Date 29(L) x 43 (H) mm

Distributed by : Morepen Bio Inc. 666, Plainsboro Road, Suite 215 Plainsboro, NJ 08536



N 3 6 2 9 9 3 0 0 0 2 4 1 6

3000 Tablets Bottle Label (NDC 62993-0002-5)

NDC 62993-0002-5

Drug Facts**Active ingredient**

(in each tablet) **Purpose**
 Docusate sodium USP 50mg Stool softener
 Sennosides 8.6 mg Laxative

Uses ■ relieves occasional constipation (irregularity)
 ■ generally produces a 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.



Docusate Sodium 50 mg & Sennosides 8.6 mg Tablets

Standardized Senna Concentrate, 8.6 mg
 Docusate Sodium, 50 mg

*Compare to active ingredient in Senokot®-S

Dual Action

Gentle, dependable constipation relief

3000 Tablets

Drug Facts (continued)

Directions ■ take preferably at bedtime or as directed by a doctor

age	Starting dosage	maximum dosage
adults and children 12 years 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 17.7 mg, sodium 4.4 mg VERY LOW SODIUM ■ store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).

Inactive ingredients colloidal silicon dioxide, croscarmellose sodium, dibasic calcium phosphate, FD&C Yellow #5 Lake*, FD&C Yellow #6 Lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, talc, titanium dioxide. *contains FD&C Yellow #5 Lake (tartrazine) as a colour additive.

Question or comments? Call toll-free 1-844-317-8549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

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No varnish Area for Lot No., Exp Date 29(L) x 43 (H) mm

Distributed by : Morepen Bio Inc. 666, Plainsboro Road, Suite 215 Plainsboro, NJ 08536



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DOCUSATE SODIUM AND SENNOSIDES

docusate sodium and sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62993-0002
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
TALC (UNII: 7SEV7J4R1U)	
POVIDONE K30 (UNII: U725QWY32X)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I3O)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Product Characteristics

Color	orange	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62993-0002-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
2	NDC:62993-0002-2	60 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
3	NDC:62993-0002-3	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
4	NDC:62993-0002-4	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
5	NDC:62993-0002-5	3000 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/01/2025	

Labeler - Morepen Laboratories Limited (650087067)

Registrant - Morepen Laboratories Limited (772653251)

Establishment

Name	Address	ID/FEI	Business Operations
Morepen Laboratories Limited		772653251	manufacture(62993-0002) , pack(62993-0002) , analysis(62993-0002) , label(62993-0002)