

CAREMOTION- sennosides tablet

US Drugs Inc.

Reference Label Set Id: c4c281e6-b449-eb75-e053-2995a90afb6f

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CareMotion Tablets
Natural Vegetable Laxative Ingredient

Active ingredient

(in each tablet)

Sen nosides 8.6 mgLaxative

Purpose

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

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. (1-800-222-1222)

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	1 /2 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 5 mg
- store at 20-25°C (68-77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

crospovidone, dichloromethane, hypromellose, isopropyl alcohol, lactose, magnesium stearate, microcrystalline cellulose, povidone

Questions or Comments? 1-866-561-2888

*This product is not manufactured or distributed by Purdue Products L.P., owner of the registered trademark Senokot

Distributed by:
 US Drugs Inc.
 11380 7th Street, Rancho Cucamonga, CA 91730, USA
 Country of Origin: India

NDC 80415-004-01



**REGULAR STRENGTH
 CARE MOTION™**

Natural Vegetable Laxative Ingredient

Gentle, Dependable
 Overnight Relief

600 Tablets



Compare to Senokot® active ingredient *



Drug Facts	
Active ingredient (in each tablet) Sennosides 8.6 mg	Purpose Laxative
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Drug Facts (continued)		
Directions ■ take preferably at bedtime or as directed by a doctor		
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Other information ■ each tablet contains: calcium 5 mg ■ store at 20 -25 °C (68 -77 °F); excursions permitted between 15°-30°C (59°-86°F)		
Inactive ingredients crospovidone, dichloromethane, hypromellose, isopropyl alcohol, lactose, magnesium stearate, microcrystalline cellulose, povidone		
Questions or Comments? 1-866-561-2888		

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THIS SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

Distributed by:
 US Drugs Inc.
 11380 7th Street, Rancho Cucamonga, CA 91730, USA
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Lot No.:
 Mfg Date:
 Exp Date:

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NDC 80415-004-02



REGULAR STRENGTH
CARE MOTION

Natural Vegetable Laxative Ingredient

Gentle, Dependable
Overnight Relief



800 Tablets
At Actual Size

Compare to Senokot® active ingredient *

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Drug Facts (continued)
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Lot No.:
Mfg Date:
Exp Date:

CAREMOTION

sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80415-004
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
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSPVIDONE (UNII: 2S7830E561)	
POVIDONE K30 (UNII: U725QWY32X)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLENE CHLORIDE (UNII: 588X2YUY0A)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80415-004-01	600 in 1 BOTTLE; Type 0: Not a Combination Product	06/14/2021	
2	NDC:80415-004-02	800 in 1 BOTTLE; Type 0: Not a Combination Product	06/14/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	06/14/2021	

Labeler - US Drugs Inc. (104282689)**Registrant** - US Drugs Inc. (104282689)

Revised: 11/2021

US Drugs Inc.