SENNA PLUS- sennosides and docusate sodium tablet
REMEDYREPACK INC.

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.

Active ingredient (in each tablet)

Docusate Sodium 50 mg
Sennosides 8.6 mg

Purpose

Stool softener

Laxative

Uses

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

Warnings

Do not use for more than 1 week unless directed by a doctor

Ask a doctor before use if you -have abdominal pain, nausea or vomiting -are taking mineral oil -have noticed a sudden change in bowel habits that lasts over 2 weeks

Stop use and ask a doctor if -you have no bowel movement within 12 hours -you have rectal bleeding. these could 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 overdose, get medical help or contact a Poison Control Center right away.

Directions

• do not exceed 8 tablets in 24 hours

Age	Starting Dose	Maximum Dose
adults and children 12 years of age and older	2 tablets once a day preferably at bedtime; increase as needed, or as directed by a doctor	4 tablets in the morning and 4 tablets at bedtime
children under 12 years	ask a doctor	

Other information

• **each tablet contains:** calcium 7 mg

• store at room temperature

Inactive ingredients

cellulose, croscarmellose sodium, dicalcium phosphate, FD and C yellow no. 5 (tartrazine), FD and C yellow no. 6, hypromellose, magnesium silicate, magnesium stearate, mineral oil, PEG, sodium benzoate, sodium lauryl sulfate, starch, stearic acid, titanium dioxide, triacetin

DRUG: Senna Plus

GENERIC: Sennosides and Docusate Sodium

DOSAGE: TABLET

ADMINSTRATION: ORAL

NDC: 70518-1590-0

COLOR: yellow

SHAPE: ROUND

SCORE: No score

SIZE: 10 mm

IMPRINT: CPC490

PACKAGING: 30 in 1 BLISTER PACK

ACTIVE INGREDIENT(S):
- DOCUSATE SODIUM 50mg in 1
- SENNOSIDES 8.6mg in 1

INACTIVE INGREDIENT(S):
- CELLULOSE, MICROCRYSTALLINE
- SODIUM BENZOATE
- FD&C YELLOW NO. 6
- SODIUM LAURYL SULFATE
- FD&C YELLOW NO. 5
- TITANIUM DIOXIDE
- STARCH, CORN
- STEARIC ACID
- CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS
- CROSCARMELLOSE SODIUM
- HYPROMELLOSES
- MAGNESIUM SILICATE
- MAGNESIUM STEARATE
- POLYETHYLENE GLYCOLS
- MINERAL OIL
- TRIACETIN

Senna Plus

Sennosides/Docusate Sodium

8.6mg/50 mg Tablet QTY: **30**

ID #: .
NDC #: 70518-1590-00
LOT #:

Expires:
Shape: Round
Ref #: 57896-0455-01

MFG: Health Star/GeriCare, Brooklyn, NY 11204

NOT FOR RETAIL SALE

Directions For Use: See Package Insert

Store at 20-25°C (68-77°F); excursions permitted to 15-30°C (59-86°F) [See USP]

Repackaged by:
RemedyRepack Inc., Indiana, PA 15701, 1-724-465-8762

SENNA PLUS

sennosides and docusate sodium tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70518-1590(NDC:57896-455)
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**
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
MAGNESIUM SILICATE (UNII: 9B9691B2N9)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	yellow	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	CPC490
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70518-1590-0	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	10/25/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	10/25/2018	

Labeler - REMEDYREPACK INC. (829572556)

Revised: 10/2019 REMEDYREPACK INC.