SENNA-PLUS- sennosides and docusate sodium tablet
Bryant Ranch Prepack

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hst 455b (555)

Active ingredient (in each tablet)

Docusate Sodium 50 mg
Sennosides 8.6 mg

Purpose

Stool softener

Laxative

Uses

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

Warnings

Do not use for more than one 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 two 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

• 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 20 mg, sodium 3 mg**
- store at controlled room temperature
  Tamper Evident: Do not use if imprinted seal under cap is missing or broken.
- product of India

Inactive ingredients

cellulose, croscarmellose sodium, dicalcium phosphate, FD&C yellow #6 lake, hypromellose, magnesium stearate, PEG, silica, talc, titanium dioxide.

HOW SUPPLIED

NDC: 71335-1674-1: 30 Tablets in a BOTTLE

NDC: 71335-1674-2: 120 Tablets in a BOTTLE

NDC: 71335-1674-3: 60 Tablets in a BOTTLE

NDC: 71335-1674-4: 90 Tablets in a BOTTLE

NDC: 71335-1674-5: 100 Tablets in a BOTTLE

NDC: 71335-1674-6: 28 Tablets in a BOTTLE

NDC: 71335-1674-7: 56 Tablets in a BOTTLE

NDC: 71335-1674-8: 14 Tablets in a BOTTLE

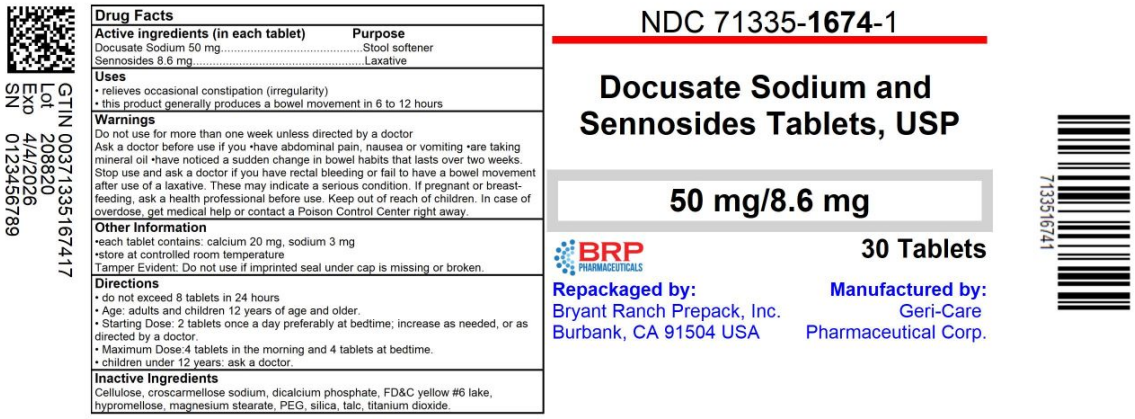
NDC: 71335-1674-9: 20 Tablets in a BOTTLE

Repackaged/Relabeled by:

Bryant Ranch Prepack, Inc.

Burbank, CA 91504

Docusate/ Sennosides 50/8.6 mg Tablet



SENNA-PLUS

sennosides and docusate sodium tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71335-1674(NDC:57896-555)
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
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TALC (UNII: 7SEV7J4R1U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	orange	**Score**	no score
Shape	ROUND	**Size**	9mm
Flavor		**Imprint Code**	PSD22
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-1674-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2021	
2	NDC:71335-1674-2	120 in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
3	NDC:71335-1674-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	09/23/2020	
4	NDC:71335-1674-4	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2024	
5	NDC:71335-1674-5	100 in 1 BOTTLE; Type 0: Not a Combination Product	08/07/2020	
6	NDC:71335-1674-6	28 in 1 BOTTLE; Type 0: Not a Combination Product	08/24/2020	
7	NDC:71335-1674-7	56 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2024	
8	NDC:71335-1674-8	14 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2024	
9	NDC:71335-1674-9	20 in 1 BOTTLE; Type 0: Not a Combination Product	04/04/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/01/2019	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-1674) , RELABEL(71335-1674)