SENNA LAXATIVE- senna tablet, film coated PD-Rx Pharmaceuticals, Inc.

Drug Facts

Active ingredient (in each tablet)

Sennosides 8.6mg

Purpose

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
- noticed a sudden change in bowel habits that last over 2 weeks

Stop use and ask a doctor if

- you have no bowel movement within 12 hours
- you have rectal bleeding

These could be 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 recommended dose

Age	Starting Dose	Maximum Dose
	preferably at bedtime;	4 tablets twice a day, in the morning and at bedtime
children under 12 years	ask a doctor	ask a doctor

Other information

- Tamper Evident: Do not use if safety seal under cap is broken or missing
- each tablet contains: calcium 20 mg, potassium 70 mg
- store at room temperature

Inactive ingredients:

croscarmellose sodium, dicalcium phosphate, hypromellose, magnesium stearate, microcrystalline cellulose, mineral oil

Questions?

Adverse drug event call: (866) 562-2756

16 HOW SUPPLIED

Senna Laxative Tablets are available as follows:

8.6 mg: Brown, round, scored tablet, debossed "SB". Available in bottles of

10 tablets (NDC 43063-838-10)

30 tablets (NDC 43063-838-30)



SENNA LAXATIVE

senna tablet, film coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43063-838(NDC:16103-363)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SENNOSIDES A AND B (UNII: 1B5FPI42EN) (SENNOSIDES A AND B - UNII:1B5FPI42EN)	SENNOSIDES A AND B	8.6 mg	

Inactive Ingredients			
Ingredient Name	Strength		
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)			
MINERAL OIL (UNII: T5L8T28FGP)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			

Product Characteristics				
Color	brown	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	S8	

Contains

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43063- 838-10	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/27/2021	
2	NDC:43063- 838-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/27/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M007	01/24/2007		

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)

Registrant - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment			
Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(43063-838)

Revised: 10/2023 PD-Rx Pharmaceuticals, Inc.