DOCUSATE SODIUM- docusate sodium capsule PD-Rx Pharmaceuticals, Inc.

Drug Facts

Active ingredient (in each softgel)

Docusate Sodium 100 mg

Purpose

Stool softener laxative

Uses

- for relief of occasional constipation (irregularity)
- this product generally produces bowel movement in 12 to 72 hours

Warnings

Do not use

if you are presently taking mineral oil, unless told to do so by a doctor.

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that last over 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 could be a signs of a serious condition.
- you need to use a laxative for more than 1 week

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 (1-800-222-1222) right away.

Directions

• take with a glass of water

years and over	take 1 to 3 softgels daily. This dose may be taken as a single daily dose or in divided dose
children 2 to under 12 years of age	take 1 softgel daily
children under 2 years	ask a doctor

Other information

- each softgel contains:sodium 5 mg
- store at 25°C (77°F); excursions permitted between 15-30°C (59-86°F)

Inactive ingredients

edible ink, FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water sorbitan, sorbitol

Questions or comments?

Call **1-877-753-3935**Monday-Friday 9AM-5PM EST

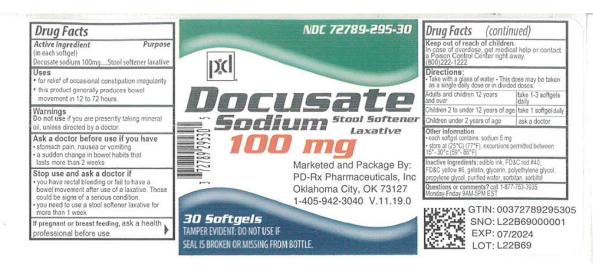
Principal Display Panel

Docusate Sodium

Stool Softener Laxative

TAMPER EVIDENT: DO NOT USE IF SEAL IS BROKEN OR MISSING FROM BOTTLE.

Product Label



DOCUSATE SODIUM

docusate sodium capsule

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72789-295(NDC:50804-862)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg	

Inactive Ingredients			
Ingredient Name	Strength		
FD&C RED NO. 40 (UNII: WZB9127XOA)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
GELATIN (UNII: 2G86QN327L)			
SORBITAN (UNII: 6O92ICV9RU)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SORBITOL (UNII: 506T60A25R)			
GLYCERIN (UNII: PDC6A3C0OX)			
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)			

Product Characteristics

Color	orange	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	P51
Contains			

ı	Packaging				
-	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:72789- 295-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/23/2022		
:	NDC:72789- 295-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/23/2022		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	05/28/2021	

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(72789-295)

Revised: 7/2024 PD-Rx Pharmaceuticals, Inc.