DOCUSATE SODIUM 50 MG- docusate sodium capsule, liquid filled Humanwell PuraCap Pharmaceutical (Wuhan), Ltd.

DOCUSATE SODIUM 50mg, Capsule, liquid filled

Drug Facts

Active ingredient (in each softgel)

Docusate Sodium 50 mg

Purpose

Stool softener

Uses

- relieves occasional constipation (irregularity)
- 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 lasts 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 signs of a serious condition.
- you need to use a stool softener 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 right away.

Directions

Take only by mouth. Doses may be taken as a single daily dose or in divided doses.

Adults and children 12 years of ages and over	take 1 to 6 softgels daily
Children 2 and under 12 years of age	take 1 to 3 softgels daily
children under 2 years of age	ask a doctor

Other information

- each softgel contains: sodium 3 mg VERY LOW SODIUM
- store at 15°-30°C (59°-86°F) Keep tightly closed.

Inactive ingredients

citric acid, D&C red #33, FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water, sorbitol special, white edible ink

Manufactured by: Humanwell PuraCap Pharmaceutical (Wuhan) Ltd. Wuhan, Hubei 430206, China

PRINCIPAL DISPLAY PANEL - Shipping Label

DOCUSATE SODIUM CAPSULES, 50 mg

Quantity : 20000 Capsules NDC. No : 53345-015-01

IMPORTANT:

Inspect immediate upon receipt. This is a bulk shipment intended for further processing only. Protect from heat, humidity, and light. Do not refrigerate.

CAUTION : "FOR FURTHER MANUFACTURING, PROCESSING OR REPACKING"

Humanwell PuraCap Pharmaceutical (Wuhan) Ltd.

No. 99, 2^{no} Shendun Road, East Lake New Technology Development District, Wuhan, Hubei 430206, P. R. China

NDC No.: 53345-015-01

Product:

DOCUSATE SODIUM CAPSULES, 50 MG

Each softgel contains: Docusate Sodium USP, 50 mg

CAUTION: FOR FURTHER MANUFACTURING, PROCESSING OR REPACKAGING

Product Code:		Quantity:		
40-00019 Lot No.: 0000000		20000 Capsules		
		Manufacturing Date:		
		00/0000		
2. This is a bulk s		diately up on receipt.		

REV-00 09/2013

DOCUSATE SODIUM	50 MG				
docusate sodium capsule, liqu	uid filled				
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (So	urce)	NDC:533	45-015
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingre	edient Name		Basis of St	rength	Strength
DOCUSATE SODIUM (UNII: F05Q2	T2JA0) (DOCUSATE - UNII:M	7P27195AG)	DOCUSATE SO	DIUM	50 mg
Inactive Ingredients					
	Ingredient Name			S	trength

FD&C RED NO.	40 (UNII: WZB9127XOA)		
D&C RED NO. 3	3 (UNII: 9DBA0SBB0L)		
FD&C YELLOW	NO. 6 (UNII: H77VEI93A8)		
GELATIN (UNII: 2	2G86QN327L)		
GLYCERIN (UNII	: PDC6A3C0OX)		
POLYETHYLEN	E GLYCOL, UNSPECIFIED (UNII: 3WJQ0	SDW1A)	
PROPYLENE GL	YCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 05	590F0K00R)		
SORBITOL (UNI			
SORBITOL (UNI			
SORBITOL (UNI	I: 506T60A25R)		
SORBITOL (UNI	I: 506T60A25R)		
SORBITOL (UNII ANHYDROUS CI	I: 506T60A25R)		
SORBITOL (UNII ANHYDROUS CI	I: 506T60A25R) ITRIC ACID (UNII: XF417D3PSL)	Score	no score
SORBITOL (UNII ANHYDROUS CI Product Cha	I: 506T60A25R) ITRIC ACID (UNII: XF417D3PSL) Aracteristics	Score Size	no score 13mm
SORBITOL (UNII ANHYDROUS CI Product Cha Color	ITRIC ACID (UNII: XF417D3PSL) aracteristics red (clear)		

Pac	kag	ing	

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53345-015- 01	1 in 1 BOX	11/12/2013	
1		20000 in 1 BAG; Type 0: Not a Combination Product		
Marketing Information				
	Marketing	Application Number or Monograph	Marketing Start	Marketing End

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	11/12/2013	

Labeler - Humanwell PuraCap Pharmaceutical (Wuhan), Ltd. (421293287)

Establishment			
Name	Address	ID/FEI	Business Operations
Humanwell PuraCap Pharmaceutical (Wuhan), Ltd.		421293287	manufacture(53345-015)

Revised: 11/2024

Humanwell PuraCap Pharmaceutical (Wuhan), Ltd.