STOOL SOFTENER LAXATIVE EXTRA STRENGTH- docusate sodium capsule, liquid filled

WALMART INC. (see also Equate)

Drug Facts

Active ingredient (in each softgel)

Docusate sodium 250 mg

Purpose

Stool softener laxative

Uses

- relieves occasional constipation
- this product generally produces a bowel movement within 12 to 72 hours

Warnings

Do not use

if you are presently taking mineral oil, unless directed 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 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 right away.

Directions

 adults and children 12 years of age and over: take 1 softgel daily or as directed by a doctor • children under 12 years of age: ask a doctor

Other information

- each softgel contains: sodium 13 mg
- store between 20-25°C(68-77°F); excursions permitted between 15-30°C (59-86°F)

Inactive Ingredients

anhydrous citric acid, FD&C red #40, FD&C yellow #6, gelatin, glycerin, isopropyl alcohol, lecithin, mannitol, mineral oil, polyethylene glycol, propylene glycol, purified water, sorbitan, sorbitol, white ink

Questions or comments?

Call 1-888-287-1915

Principal Display Panel

EXTRA STRENGTH

Stool Softener

Docusate sodium 250 mg

Stool Softener Laxative

- Stimulant-free
- Relief of occasional constipation

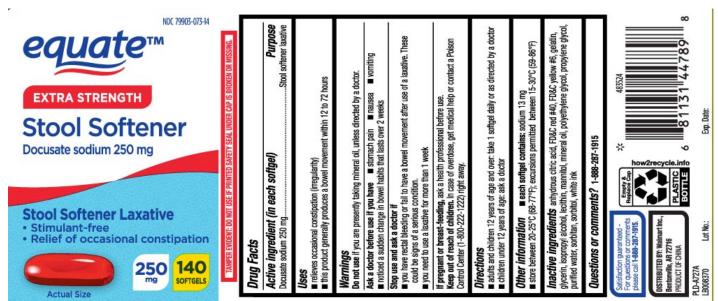
SOFTGELS

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

Distributed BY: Walmart Inc.,

Bentonville, AR 72716

Product Label



EQUATE Extra Strength Stool Softener

STOOL SOFTENER LAXATIVE EXTRA STRENGTH

docusate sodium capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-073
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	250 mg	

Inactive Ingredients	
Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MANNITOL (UNII: 30WL53L36A)	
MINERAL OIL (UNII: T5L8T28FGP)	
SORBITAN (UNII: 6O92ICV9RU)	
MANNITOL (UNII: 30WL53L36A) MINERAL OIL (UNII: T5L8T28FGP)	

Product Characteristics			
Color	orange	Score	no score
Shape	CAPSULE	Size	20mm
Flavor		Imprint Code	P4
Contains			

Packaging				
# Item (Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:7990	03-073- 140 in Produc	1 BOTTLE; Type 0: Not a Combination t	06/25/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	06/25/2021	

Labeler - WALMART INC. (see also Equate) (051957769)

Revised: 6/2024 WALMART INC. (see also Equate)