

STOOL SOFTENER AND STIMULANT LAXATIVE- docusate sodium and sennosides tablet
TARGET CORPORATION

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

209R TARGET SENNOSIDES AND DOC SOD TABS 8.6/50 MG NDC 11673-209-01

ACTIVE INGREDIENTS

active ingredients (each tablet)

Docusate Sodium 50 mg

Sennosides 8.6 mg

INACTIVE INGREDIENTS

CARNAUBA WAX, COLLOIDAL SILICON DIOXIDE, CROSCARMELLOSE SODIUM, DIBASIC CALCIUM PHOSPHATE DIHYDRATE, FD-C RED NO. 40 ALUMINUM LAKE, HYPROMELLOSE, MAGNESIUM STEARATE, MICROCRYSTALLINE CELLULOSE, POLYETHYLENE GLYCOL, SODIUM BENZOATE, STEARIC ACID, TITANIUM DIOXIDE

PURPOSE

Stool Softener

Stimulant Laxative

INDICATIONS AND USAGE

RELIEVES OCCASIONAL CONSTIPATION

GENERALLY PRODUCES BOWEL MOVEMENT IN 6-12 HOURS

WARNINGS

DO NOT USE

LAXATIVE PRODUCTS FOR LONGER THAN 1 WEEK UNLESS DIRECTED BY A DOCTOR

IF YOU ARE PRESENTLY TAKING MINERAL OIL UNLESS DIRECTED BY A DOCTOR

DOSAGE AND ADMINISTRATION

ADULTS AND CHILDREN 12 YEARS AND OVER: 2-4 TABLETS ONCE DAILY OR IN DIVIDED DOSES

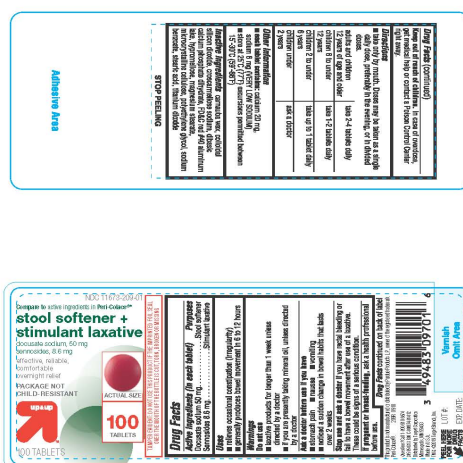
CHILDREN 6 TO UNDER 12 YEARS: 1-2 TABLETS ONCE DAILY OR IN DIVIDED DOSES

CHILDREN 2 TO UNDER 6 YEARS; 1/2 -1 TABLET ONCE DAILY OR IN DIVIDED DOSES

CHILDREN UNDER 2 YEARS: ASK A DOCTOR

KEEP OUT OF REACH OF CHILDREN

PRINCIPAL DISPLAY PANEL



STOOL SOFTENER AND STIMULANT LAXATIVE

docusate sodium and sennosides tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-209
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
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	10 mm
Flavor		Imprint Code	TCL097
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-209-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	04/10/2019	

Labeler - TARGET CORPORATION (006961700)**Registrant** - TIME CAP LABORATORIES, INC. (037052099)**Establishment**

Name	Address	ID/FEI	Business Operations
TIME CAP LABORATORIES, INC.		037052099	manufacture(11673-209)

