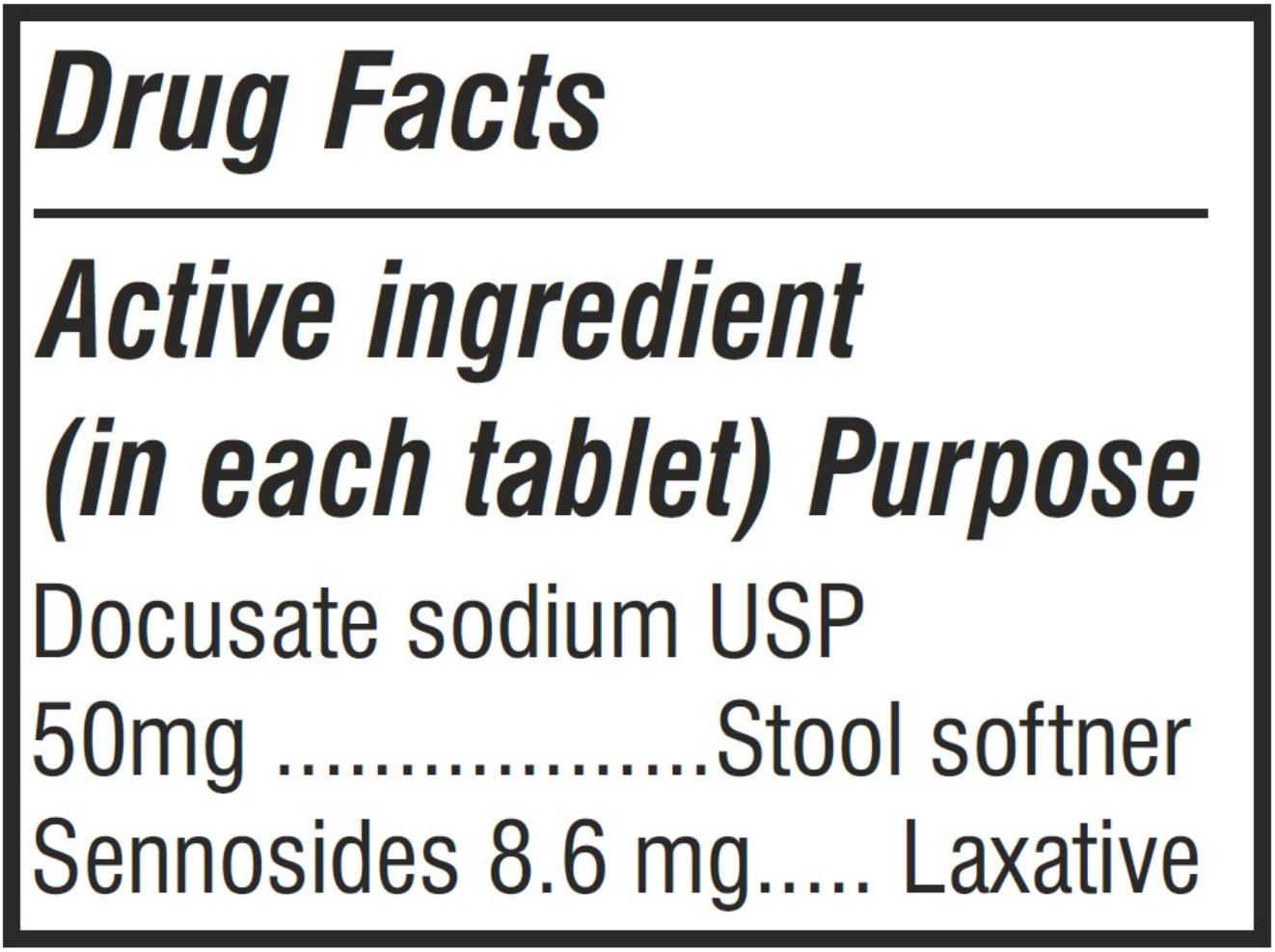
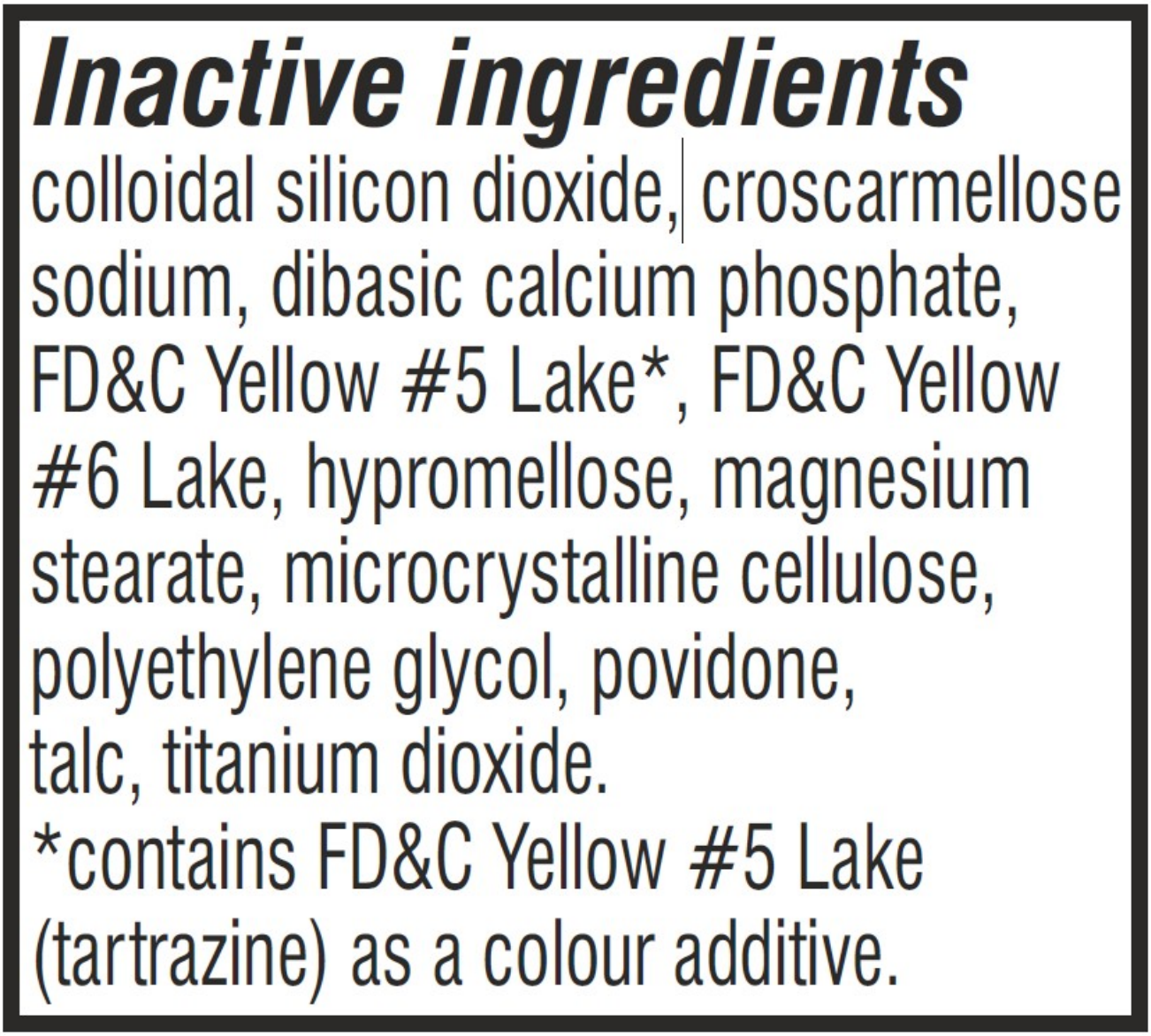
DOCUSATE SODIUM AND SENNOSIDES- docusate sodium and sennosides tablet
Morepen Laboratories Limited

Docusate Sodium 50 mg and Sennosides 8.6 mg Tablets

Active Ingredients Label



Inactive Ingredients Section Label



Purpose Section Label

Drug Facts

Active ingredient (in each tablet) Purpose

Docusate sodium USP
50mg .....Stool softner
Sennosides 8.6 mg..... Laxative

Indications & Usage Seciton Label

Uses

- relieves occasional constipation (irregularity)
- generally produces a bowel movement in 6-12 hours

Dosage & Administration Section Label

Directions

■ take preferably at bedtime or as directed by a doctor

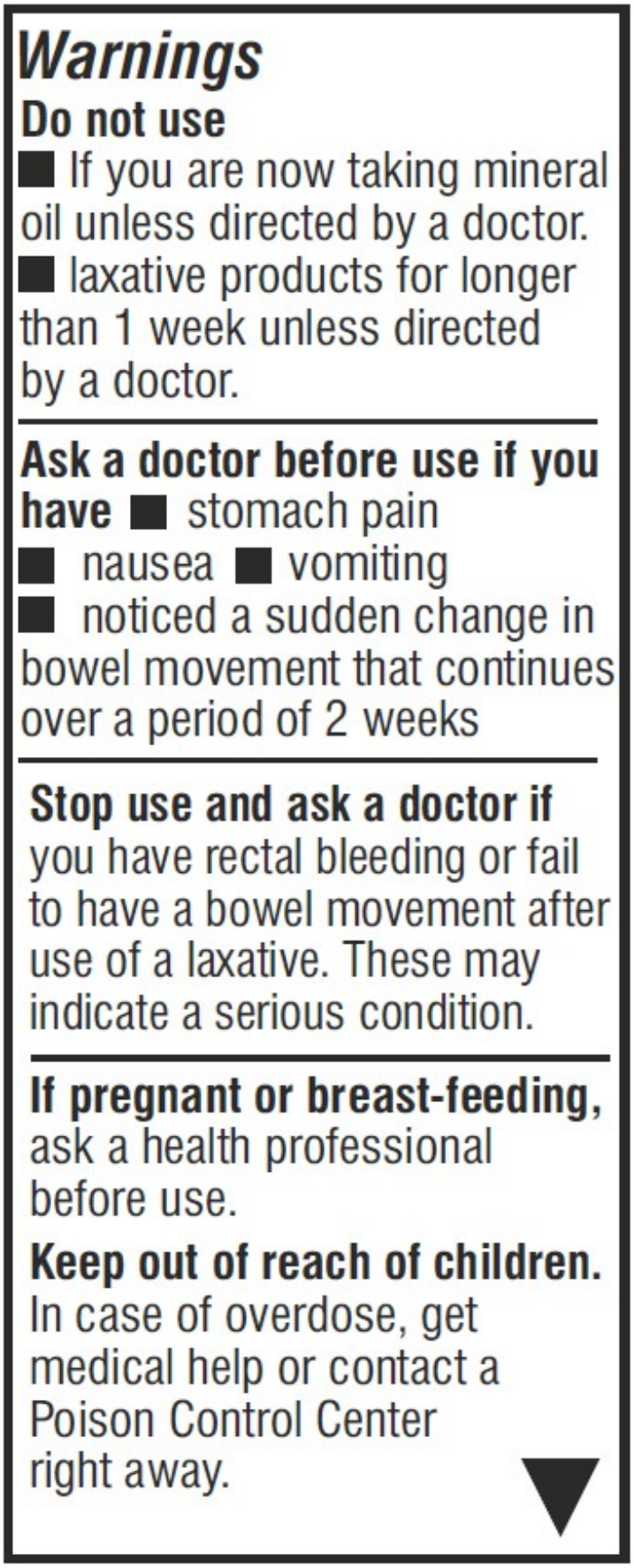
age	Starting dosage	maximum dosage
adults and children 12 years and over	2 tablets once a day	4 tablets twice a day
children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
children 2 to under 6 years	½ tablet once a day	1 tablet twice a day
children under 2 years	ask a doctor	ask a doctor

Questions Section Label

Question or comments?

Call toll- free 1-844-317-6549 Mon-Fri 9AM-5PM EST or contact the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

Warnings Section Label



Keep out or Reach of Children Label

Keep out of reach of children.
In case of overdose, get medical help or contact a Poison Control Center right away.

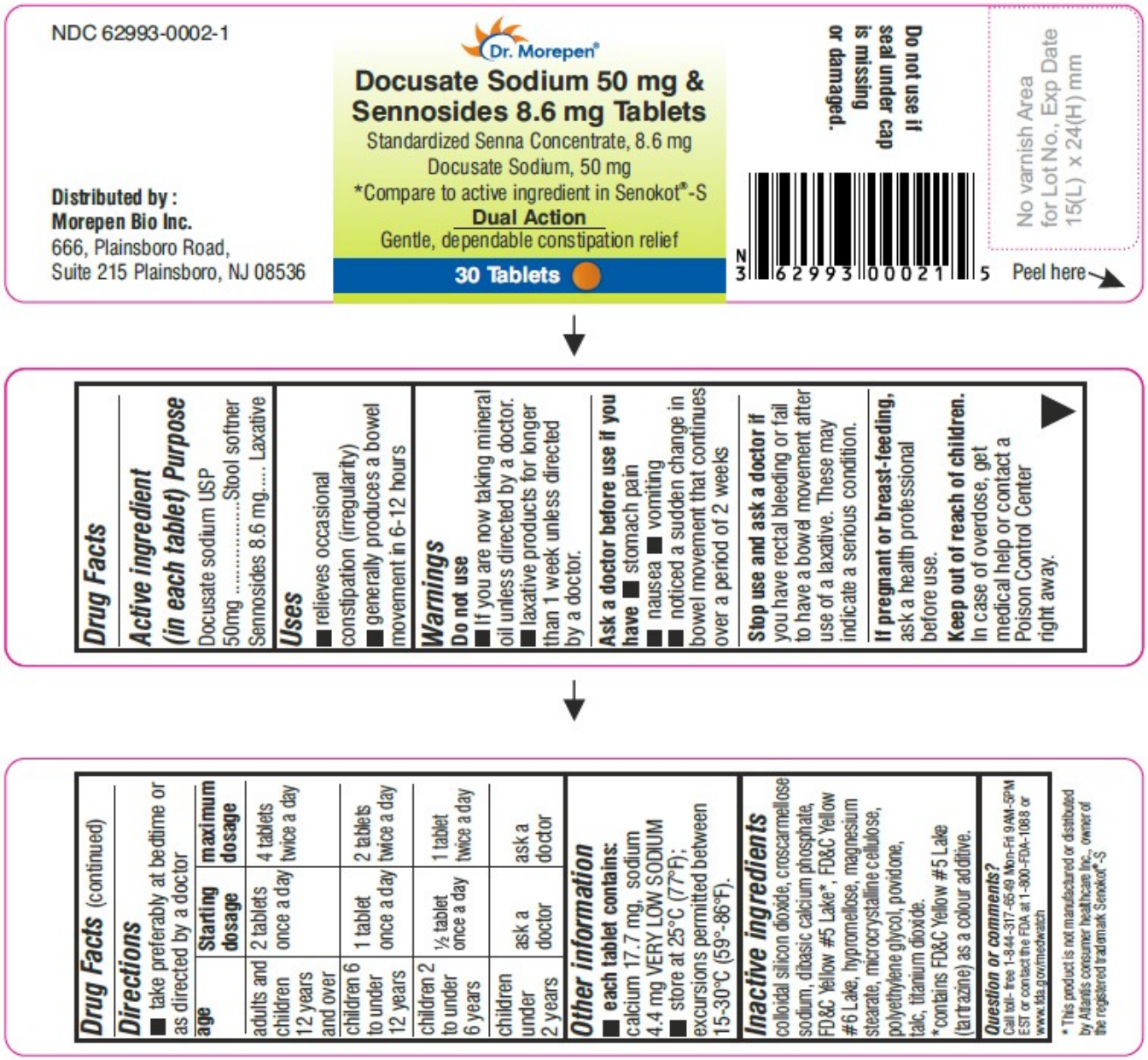
Storage and Handling Section Label

Other information

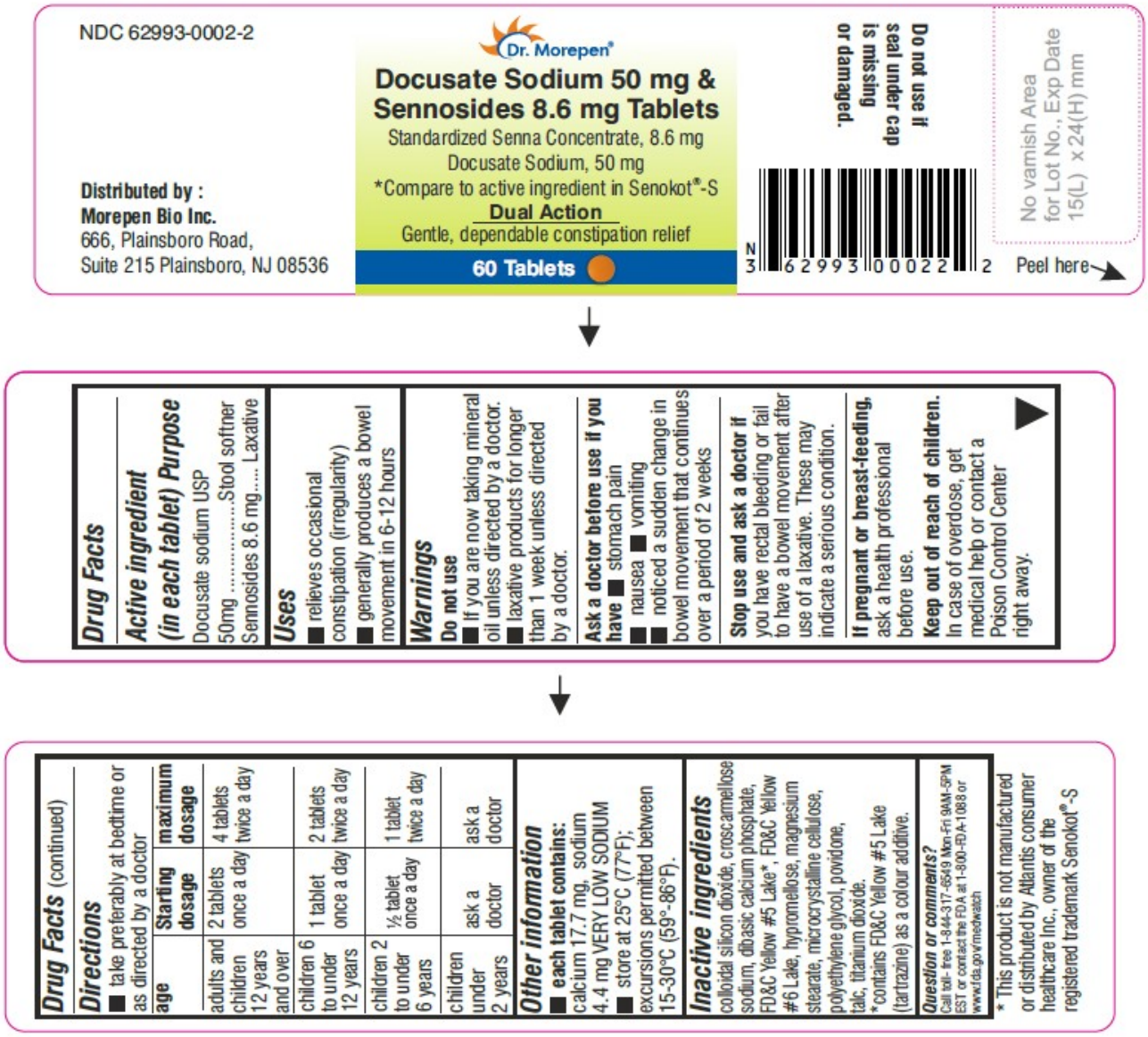
■ **each tablet contains:**
calcium 17.7 mg, sodium 4.4 mg VERY LOW SODIUM

■ store at 25°C (77°F); excursions permitted between 15-30°C (59°-86°F).

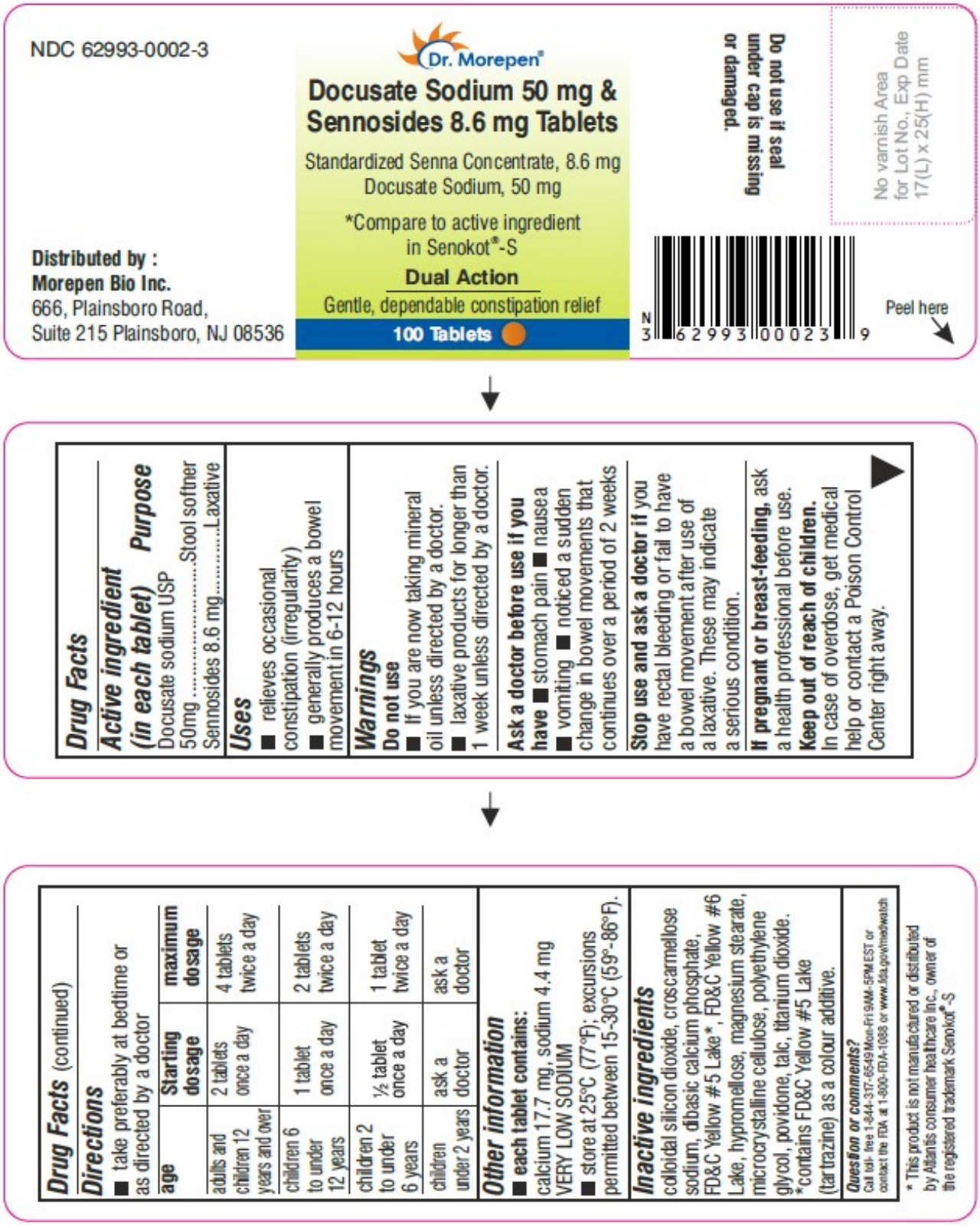
30 Tablets Bottle Label (NDC 62993-0002-1)



60 Tablets Bottle Label (NDC 62993-0002-2)



100 Tablets Bottle Label (NDC 62993-0002-3)



1000 Tablets Bottle Label (NDC 62993-0002-4)

3000 Tablets Bottle Label (NDC 62993-0002-5)

DOCUSATE SODIUM AND SENNOSIDES

docusate sodium and sennosides tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62993-0002
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	50 mg
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)	SENNOSIDES	8.6 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CALCIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: L11K75P92J)	
TALC (UNII: 7SEV7J4R1U)	
POVIDONE K30 (UNII: U725QWY32X)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Product Characteristics

Color	orange	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62993-0002-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
2	NDC:62993-0002-2	60 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
3	NDC:62993-0002-3	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
4	NDC:62993-0002-4	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	
5	NDC:62993-0002-5	3000 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	10/01/2025	

Labeler - Morepen Laboratories Limited (650087067)

Registrant - Morepen Laboratories Limited (772653251)

Establishment

Name	Address	ID/FEI	Business Operations
Morepen Laboratories Limited		772653251	manufacture(62993-0002) , pack(62993-0002) , analysis(62993-0002) , label(62993-0002)

Revised: 10/2025 Morepen Laboratories Limited