

SENNOSIDES, DOCUSATE SODIUM- sennosides, docusate sodium tablet, film coated

Major Pharmaceuticals

3052R - Major

Drug Facts

Active ingredients (in each tablet)

Docusate sodium 50 mg

Sennosides 8.6 mg

Purposes

Stool softener

Stimulant laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 6 to 12 hours
- laxative products for longer than 1 week unless told to do so by a doctor
- 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 two 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.

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, preferably in the evening, or in divided doses.

| | |
|---------------------------------------|---------------------------|
| adults and children 12 years and over | take 2-4 tablets daily |
| children 6 to under 12 years of age | take 1-2 tablets daily |
| children 2 to under 6 years of age | take up to 1 tablet daily |
| children under 2 years | ask a doctor |

- **each tablet contains:** calcium 5 mg
- Store in a dry place at 15° – 30°C (59° – 86°F).

corn starch, croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #1 lake, FD&C red #40 lake, FD&C yellow #6 lake, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, mineral oil, polyethylene glycol, sodium benzoate, sodium lauryl sulfate, stearic acid, talc, titanium dioxide.

Questions or comments? 1-800-231-4670

Keep this carton for complete product information

Tamper Evident:

Do not use if sealed blister units are broken or damaged.

Distributed by:

MAJOR® PHARMACEUTICALS

Indianapolis, IN 46268

(800) 616-2471

www.majorpharmaceuticals.com

Major

NDC 0904-7440-61

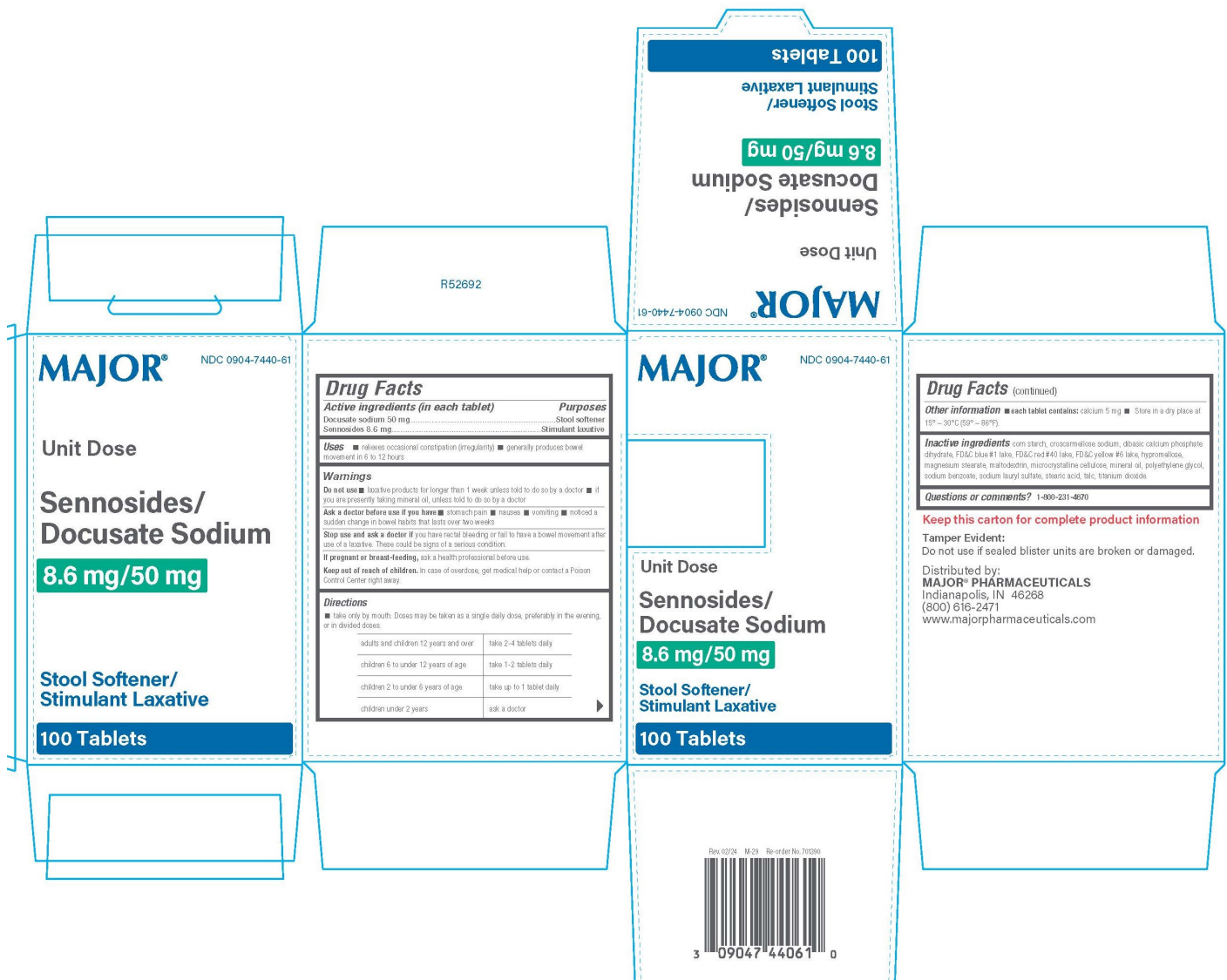
Unit Dose

Sennosides/Docusate Sodium

8.6mg/50mg

Stool Softener/Stimulant Laxative

100 Tablets



SENNOSIDES, DOCUSATE SODIUM

sennosides, docusate sodium tablet, film coated

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0904-7440 |
| 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 |
|---|----------|
| MALTODEXTRIN (UNII: 7CVR7L4A2D) | |
| CROSCARMELLOSE SODIUM (UNII: M28OL1HH48) | |

| |
|---|
| STARCH, CORN (UNII: O8232NY3SJ) |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J) |
| MAGNESIUM STEARATE (UNII: 70097M6130) |
| POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) |
| STEARIC ACID (UNII: 4ELV7Z65AP) |
| DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP) |
| FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q) |
| HYPROMELLOSES (UNII: 3NXW29V3WO) |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) |
| FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM) |
| FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T) |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) |
| SODIUM BENZOATE (UNII: OJ245FE5EU) |
| TALC (UNII: 7SEV7J4R1U) |
| LIGHT MINERAL OIL (UNII: N6K5787QVP) |

Product Characteristics

| | | | |
|-----------------|-------|---------------------|----------|
| Color | red | Score | no score |
| Shape | ROUND | Size | 10mm |
| Flavor | | Imprint Code | 49;0 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0904-7440-61 | 10 in 1 BOX, UNIT-DOSE | 05/01/2024 | |
| 1 | | 10 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M007 | 05/01/2024 | |

Labeler - Major Pharmaceuticals (191427277)

Revised: 5/2024

Major Pharmaceuticals