# STOOL SOFTENER REGULAR STRENGTH- docusate sodium capsule, liquid filled Spirit Pharmaceuticals LLC

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#### Stool Softener Regular Strength

#### **Drug Facts**

#### Active Ingredient (in each tablet)

Docusate sodium 100 mg

#### **Purpose**

Stool softener

#### Uses

• relieves occasional constipation (irregularity) • generally produces bowel movement in 12–72 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

fever • mucus in the stool

# Ask a doctor or pharmacist before use if you are 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. (1-800-222-1222)

#### **Directions**

take only by mouth. Doses may be taken as asingle daily dose or in divided doses.

| adults & children 12 years of age & over | take 1-3 softgels daily |
|--|-------------------------|
| children 2 to under 12 years of age      | take 1 softgel daily    |
| children under 2 years of age            | ask a doctor            |

#### Other information

• store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) • Keep tightly closed

### Inactive ingredients

FD&C Red #40, FD&C Yellow #6, gelatin, glycerin, polyethylene glycol-400, povidone, propylene glycol, purified water, sorbitol solution

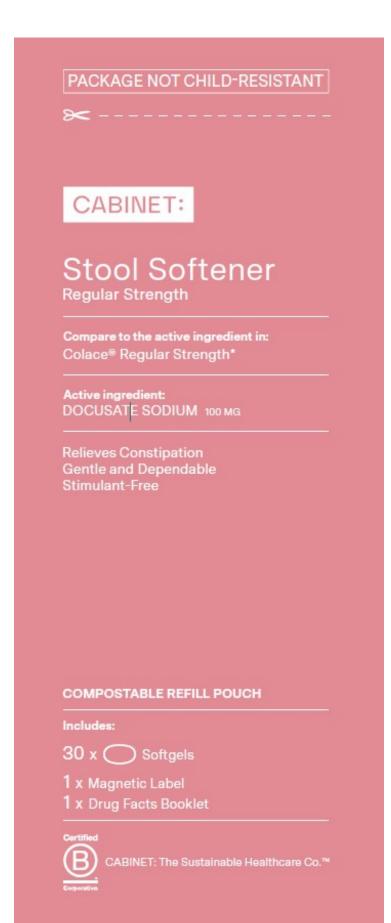
# Questions or comments?

1-888-333-9792

# Distributed by:

Cabinet Health P.B.C.

#### **Pouch**



#### STOOL SOFTENER REGULAR STRENGTH

docusate sodium capsule, liquid filled

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68210-4176 Route of Administration ORAL

| Active Ingredient/Active Moiety                                 |                          |          |  |  |
|---|--------------------------|----------|--|--|
| Ingredient Name   | <b>Basis of Strength</b> | Strength |  |  |
| DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG) | DOCUSATE SODIUM          | 100 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 400 (UNII: B697894SGQ) |          |  |  |
| POVIDONE (UNII: FZ989GH94E)                |          |  |  |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)        |          |  |  |
| WATER (UNII: 059QF0KO0R)                   |          |  |  |
| SORBITOL SOLUTION (UNII: 8KW3E207O2)       |          |  |  |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)        |          |  |  |

| Product Characteristics |      |              |          |  |
|-------------------------|------|--------------|----------|--|
| Color                   | red  | Score        | no score |  |
| Shape                   | OVAL | Size         | 12mm     |  |
| Flavor                  |      | Imprint Code | 125      |  |
| Contains                |      |              |          |  |

| l | Packaging                       |                      |  |                       |  |
|---|---------------------------------|----------------------|--|-----------------------|--|
|   | # Item Code Package Description |                      | Marketing Start<br>Date                          | Marketing End<br>Date |  |
|   |                                 | NDC:68210-<br>4176-3 | 30 in 1 POUCH; Type 0: Not a Combination Product | 11/22/2021            |  |

| Marketing Information                                       |      |                         |                       |
|---|------|-------------------------|-----------------------|
| Marketing Application Number or Monograph Category Citation |      | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug  | M007 | 11/22/2021              |                       |
|   |      |                         |                       |

# **Labeler -** Spirit Pharmaceuticals LLC (179621011)

Revised: 12/2024 Spirit Pharmaceuticals LLC