

**STOOL SOFTENER WITH STIMULANT LAXATIVE- docusate sodium and sennosides tablet**  
**CVS Pharmacy**

*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.*

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**Drug Facts**

**Active ingredients (in each tablet)**

Docusate sodium 50 mg

Sennosides 8.6 mg

**Purpose**

Stool softener

Stimulant laxative

**Uses**

- for overnight relief from occasional constipation (irregularity)
- generally produces bowel movement in 6 to 12 hours

**Warnings**

**Do not use**

- laxative products for longer than 1 week unless directed by a doctor
- if you are now 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 lasts over a period of 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 may indicate 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 older | 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                       | ask a doctor              |

### Other information

- **each tablet contains:** calcium 30 mg
- **each tablet contains:** sodium 6 mg VERY LOW SODIUM
- store at 25°C (77°F); excursions permitted between 15-30°C(59-86°F)

### Inactive ingredients

carnauba wax\*, croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol\*, silicon dioxide, sodium benzoate\*, stearic acid, talc\*, titanium dioxide

\*contains one or more of these ingredients

### Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

### Principal Display Panel

Compare to the active ingredients in Colace® 2-IN1†

Stool Softener

DOCUSATE SODIUM 50 mg &

SENNOSIDES 8.6 mg

Plus stimulant laxative

Convenient and predictable overnight relief of constipation

TABLETS

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS**

BROKEN OR MISSING.

†This product is not manufactured or distributed by Avrio Health L.P., distributor of Colace® 2-IN-1.

Distributed by: CVS Pharmacy, Inc.  
One CVS Drive, Woonsocket, RI 02895

Product Label

VALUE SIZE 400 Tablets

CVSHealth

Compare to the active ingredients in Colace® 2-IN-1\* NDC 69842-829-40

Stool Softener

DOCUSATE SODIUM 50 mg  
SENNOSIDES 8.6 mg

Plus stimulant laxative

Convenient and predictable overnight relief of constipation

Actual Size

400 TABLETS

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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.

Drug Facts (continued under label)

This product is not manufactured or distributed by Avrio Health L.P., distributor of Colace® 2-IN-1.

Distributed by: CVS Pharmacy, Inc.  
One CVS Drive, Woonsocket, RI 02895  
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Exp. Date:

CVS Quality

Money Back Guarantee

PLD-ASU  
LB008825

Lot No.:

PEEL HERE

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CVS HEALTH Stool Softener plus Stimulant Laxative

STOOL SOFTENER WITH STIMULANT LAXATIVE

docusate sodium and sennosides tablet

| Product Information             |                |                    |               |
|---------------------------------|----------------|--------------------|---------------|
| Product Type                    | HUMAN OTC DRUG | Item Code (Source) | NDC:69842-829 |
| 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             |                    |
| CARNAUBA WAX (UNII: R12CBM0EIZ)                                 |                  |                                                             |                      |                    |
| CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)                        |                  |                                                             |                      |                    |
| DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)          |                  |                                                             |                      |                    |
| FD&C RED NO. 40 (UNII: WZB9127XOA)                              |                  |                                                             |                      |                    |
| HYPROMELLOSES (UNII: 3NXW29V3WO)                                |                  |                                                             |                      |                    |
| MAGNESIUM STEARATE (UNII: 70097M6I30)                           |                  |                                                             |                      |                    |
| CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)                  |                  |                                                             |                      |                    |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)             |                  |                                                             |                      |                    |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4)                              |                  |                                                             |                      |                    |
| SODIUM BENZOATE (UNII: OJ245FE5EU)                              |                  |                                                             |                      |                    |
| STEARIC ACID (UNII: 4ELV7Z65AP)                                 |                  |                                                             |                      |                    |
| TALC (UNII: 7SEV7J4R1U)                                         |                  |                                                             |                      |                    |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                             |                  |                                                             |                      |                    |
| ALUMINUM OXIDE (UNII: LMI26O6933)                               |                  |                                                             |                      |                    |
| FD&C BLUE NO. 2 (UNII: L06K8R7DQK)                              |                  |                                                             |                      |                    |
|                                                                 |                  |                                                             |                      |                    |
| Product Characteristics                                         |                  |                                                             |                      |                    |
| Color                                                           | red              | Score                                                       | no score             |                    |
| Shape                                                           | ROUND            | Size                                                        | 10mm                 |                    |
| Flavor                                                          |                  | Imprint Code                                                | TCL097;PSD21;S44     |                    |
| Contains                                                        |                  |                                                             |                      |                    |
|                                                                 |                  |                                                             |                      |                    |
| Packaging                                                       |                  |                                                             |                      |                    |
| #                                                               | Item Code        | Package Description                                         | Marketing Start Date | Marketing End Date |
| 1                                                               | NDC:69842-829-10 | 100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/30/2019           | 04/30/2025         |
| 2                                                               | NDC:69842-829-40 | 400 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/30/2019           | 04/30/2025         |
| 3                                                               | NDC:69842-829-25 | 250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/30/2019           | 04/30/2025         |
|                                                                 |                  |                                                             |                      |                    |
| Marketing Information                                           |                  |                                                             |                      |                    |
| Marketing Category                                              |                  | Application Number or Monograph Citation                    | Marketing Start Date | Marketing End Date |
| OTC monograph not final                                         |                  | part334                                                     | 04/30/2019           | 04/30/2025         |

