DOCUSATE SODIUM- docusate sodium capsule, liquid filled
NCS HealthCare of KY, LLC dba Vangard Labs

Padagis Docusate Sodium 100 mg Drug Facts

Active ingredient (in each softgel)

Docusate sodium 100 mg

Purpose

Stool Softener Laxative

Uses

- relieves occasional constipation (irregularity)
- generally produces bowel movement in 12 to 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

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that lasts over 2 weeks

Ask a doctor or pharmacist before use if you are

taking any other drug. Take this product two or more hours before or after other drugs. Laxatives may affect how other drugs work.

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 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 a single daily dose or in divided doses.

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

Other information

- **each softgel contains**: sodium 6 mg
- very low sodium
- store at 20-25°C (68-77°F)

Inactive ingredients

D&C red no. 33, edible ink, FD&C blue no. 1, FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, polyethylene glycol, propylene glycol, purified water, sorbitol sorbitan solution, titanium dioxide

Questions or comments?

1-800-719-9260

Principal Display Panel - BINGO CARD

VG NDC 0615-8041-39

Docusate Sodium
Softgels 100 mg

LOT 8041-

EXP

QTY 30

4580248678 39

Stool
Softener
Laxative

(Padagis NDC 45802-486-78)

Docusate Sodium
Softgels 100 mg

TAMPER EVIDENT

8041-AA-39 v04

Mfg By Patheon for
Padagis
(NDC 45802-486-78)
PKG BY VANGARD
GLASGOW, KY 42141

		16	8
30	23	15	7
29	22	14	6
28	21	13	5
27	20	12	4
26	19	11	3
25	18	10	2
24	17	9	1

Omnicare **The overall configuration of this package is a trademark of Omnicare, Inc.**

Received: _____

	24_____	16_____	8_____
31_____	23_____	15_____	7_____
30_____	22_____	14_____	6_____
29_____	21_____	13_____	5_____
28_____	20_____	12_____	4_____
27_____	19_____	11_____	3_____
26_____	18_____	10_____	2_____
25_____	17_____	9_____	1_____

Start Date _____ Start Time_____

See Bottle Label for Drug Facts
and full labeling information.

Store at 20°-25°C (68°-77°F)

FOR INSTITUTIONAL USE ONLY

Pkg by Vangard, Glasgow, KY 42141 Docusate Sodium Softgel 100 mg LOT 8041- 4580248678	Pkg by Vangard, Glasgow, KY 42141 Docusate Sodium Softgel 100 mg LOT 8041- 4580248678	Pkg by Vangard, Glasgow, KY 42141 Docusate Sodium Softgel 100 mg LOT 8041- 4580248678	8041-AA-B-v04 Vangard Labs Glasgow, KY 42141 LOT 8041- 4580248678
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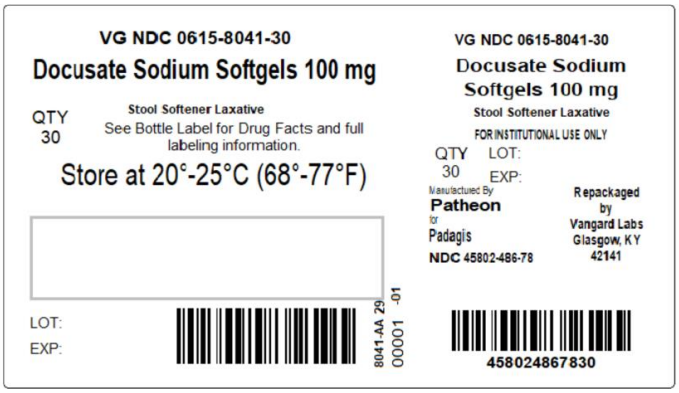
Principal Display Panel - UNIT DOSE



DOCUSATE SODIUM

docusate sodium capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0615-8041(NDC:45802-486)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color	red, white (to off beige)	**Score**	no score
Shape	OVAL (softgel)	**Size**	13mm
Flavor		**Imprint Code**	L486
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0615-8041-39	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/14/2016	
2	NDC:0615-8041-30	6 in 1 BOX, UNIT-DOSE	04/14/2016	
2		5 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	505G(a)(3)	10/28/2008	

Labeler - NCS HealthCare of KY, LLC dba Vangard Labs (050052943)

Establishment

Name	Address	ID/FEI	Business Operations
NCS HealthCare of KY, LLC dba Vangard Labs		050052943	repack(0615-8041)