GOOD NEIGHBOR PHARMACY GENTLE LAXATIVE- bisacodyl suppository
AmerisourceBergen Drug Corp

DRUG FACTS

Active ingredient

Bisacodyl USP, 10 mg

Purpose

Stimulant Laxative

Uses

- For relief of occasional constipation and irregularity
- -This product generally produces bowel movement in 15 minutes to 1 hour

Warnings

For rectal use only.

- stomach pain, nausea or vomiting
- noticed a sudden change in bowel habits that persists over a period of two weeks

When using this product

May cause abdominal discomfort, faintness, rectal burning, and mild cramps

Stop use and ask a doctor if

- if you have rectal bleeding or fail to have bowel movement after using a laxative. This may indicate 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

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 12 years of age and older	Children 6 to under 12 years	Children under 6
One suppository once daily	1/2 suppository once	

daily Ask doctor.

-Detach one suppository from the strip and remove from foil - Carefully insert one suppositry well into the rectum

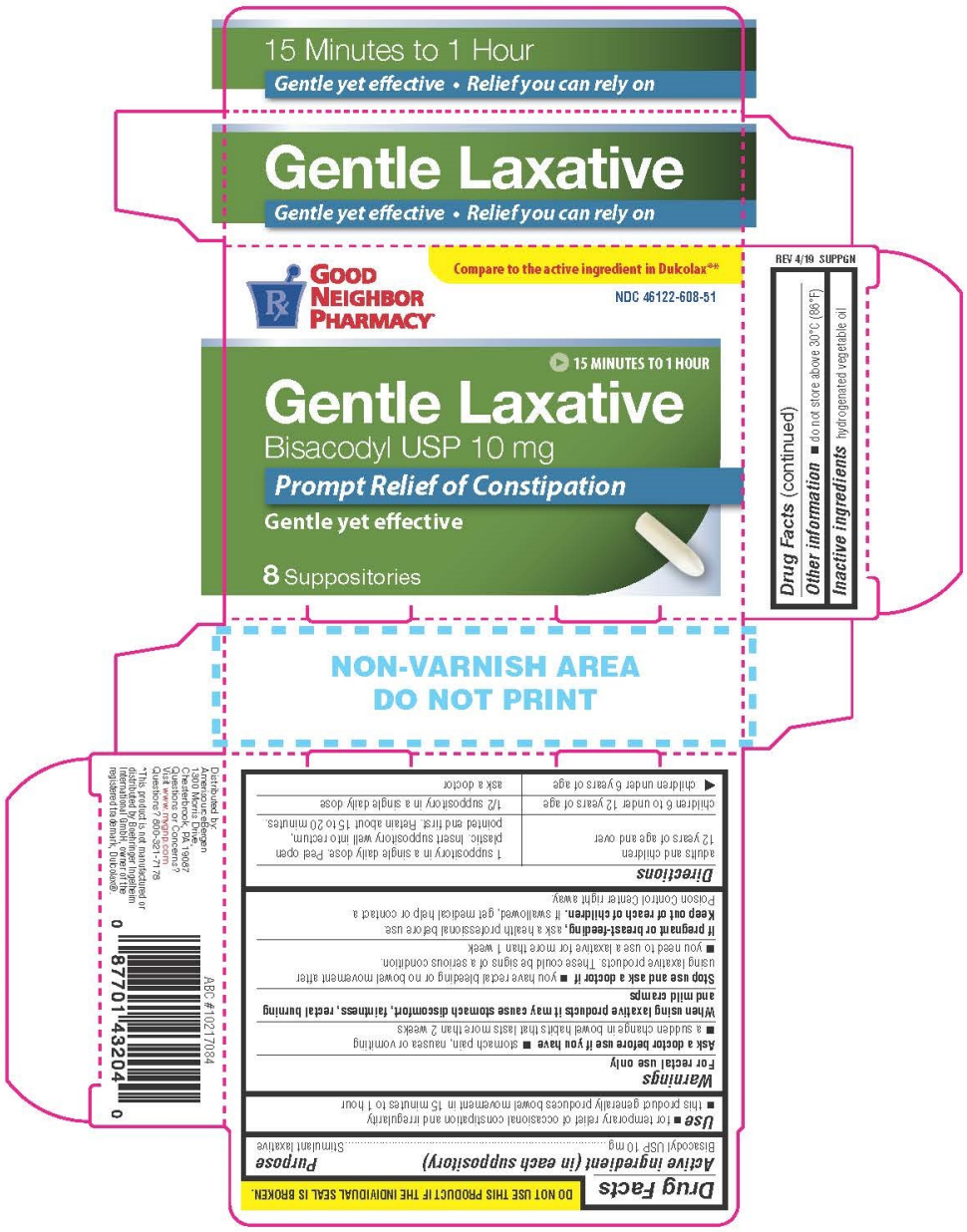
-Do not use more than once per day

Other Information

- do not store above 30°C (86°F)

Inactive Ingredients

hydrogenated vegetable oil



GOOD NEIGHBOR PHARMACY GENTLE LAXATIVE

bisacodyl suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:46122-608
Route of Administration	RECTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISACODYL (UNII: 10X0709Y6I) (DEACETYLBISACODYL - UNII:R09078E41Y)	BISACODYL	10 mg in 2000 mg

Inactive Ingredients

Ingredient Name	Strength
FAT, HARD (UNII: 8334LX7S21)	

Product Characteristics

Color	white	**Score**	
Shape	BULLET	**Size**	34mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-608-51	2 in 1 CARTON	06/26/2019	
1		40 mg 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	06/26/2019	

Labeler - AmerisourceBergen Drug Corp (007914906)

Registrant - Reese Pharmaceutical Co (004172052)

Establishment

Name	Address	ID/FEI	Business Operations
DSC Laboratories, Inc.		097807374	manufacture(46122-608)

Revised: 12/2024 AmerisourceBergen Drug Corp