ONELAX- bisacodyl suppository
AKRON PHARMA INC

OneLAX
BISACODYL 10 mg Suppositories
Fast Acting Stimulant Laxative

Active Ingredients (per suppository)

Bisacodyl USP, 10 mg

Purpose

Laxative

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Uses

- for relief of occasional constipation
- this product usually produces bowel movement in 1/4 to 1 hour

Warnings

For rectal use only.

Do not use laxative products

- when abdominal pain, nausea or vomiting are present
- for a period longer than one week

Ask a doctor before use if you have

noticed a sudden change in bowel habits that persist over a period of 2 weeks

When using this product

it may cause abdominal discomfort, faintness, rectal burning and mild cramps

Stop use and ask a doctor if

rectal bleeding occurs, or you fail to have a bowel movement after using a laxative. This may indicate a serious condition.

If pregnant or breast-feeding

ask a health professional before use.

Directions

- adults and children 12 years of age and older
- detach one suppository from the strip
- remove from wrapper before inserting into the rectum
- the rectal suppository dose is one suppository per day or as directed by a doctor
- children under 12 years of age consult a doctor

Other information

store below 30° C (86° F)

Inactive ingredient

hydrogenated vegetable oil

Manufactured for:
Akron Pharma Inc.,
Fairfield, NJ-07004
www.akronpharma.com

OneLAX™

NDC 71399-8460-2

Compare to the active ingredient in Dulcolax® Suppositories*

Bisacodyl 10mg Suppositories

Fast Acting Stimulant Laxative

12 SUPPOSITORIES

TAMPER EVIDENT: SUPPOSITORIES ARE WRAPPED IN TAMPER EVIDENT SEALED WRAPPERS. DO NOT USE IF WRAPPER IS TORN OR OPEN WHEN PURCHASED.

Bisacodyl 10mg Suppositories

Drug Facts

Active ingredient (in each suppository) — **Purpose**

Bisacodyl USP 10 mg ..... Stimulant laxative

Uses

- relieves occasional constipation
- this product generally produces a bowel movement in 15 minutes to 1 hour

Drug Facts (continued)

Warnings For rectal use only

Do not use when abdominal pain, nausea, or vomiting are present.

Ask a doctor before use if you have a sudden change in bowel habits that lasts over a period of 2 weeks.

When using this product you may experience ■ rectal burning ■ mild cramps ■ abdominal discomfort ■ faintness

Stop use and ask a doctor if ■ you have rectal bleeding ■ you fail to have a bowel movement after using a laxative. These may be signs of a serious condition.

■ you need to use a laxative for more than one week

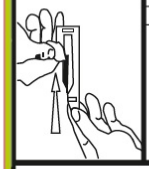
If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- tear off one suppository from the strip
- tear and peel the wrapper in a downward motion
- continue tearing downward to almost the full length of the suppository ***(see illustration at the right)***
- gently remove the suppository from the wrapper
- lie on your side and push suppository, with pointed end first, high into the rectum so it will not slip out
- retain it for 15 to 20 minutes
- **Do not exceed recommended dose** ■ if suppository seems soft, place in refrigerator for a short time before use

Adults and children 12 years and over	1 suppository once daily
Children under 12 years	Ask a doctor



Other information ■ store at temperatures below 30°C (86°F)

Inactive ingredient hydrogenated vegetable oil

Akron Pharma

Manufactured for: Akron Pharma Inc., Fairfield, NJ-07004

www.akronpharma.com

Questions or comments? toll-free (877)225-6999

*This product is not manufactured or distributed by Boehringer Ingelheim Pharmaceuticals, owner of the registered trademark Dulcolax®

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ONELAX

bisacodyl suppository

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-8460
Route of Administration	RECTAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Product Characteristics

Color	white (WHITE TO OFF WHITE)	**Score**	
Shape	OVAL (TORPEDO SHAPE)	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-8460-2	12 in 1 BOX	09/13/2024	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:71399-8460-5	50 in 1 BOX	09/13/2024	
2		1 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	09/13/2024	

Labeler - AKRON PHARMA INC (067878881)

Revised: 9/2024 AKRON PHARMA INC