

**CLEAR LAX- polyethylene glycol 3350 powder, for solution
H E B**

HEB Clearlax® Drug Facts

Active ingredient (in each dose)

Polyethylene Glycol 3350, 17 g (cap filled to line)

Purpose

Osmotic Laxative

Use

- relieves occasional constipation (irregularity)
- generally produces a bowel movement in 1 to 3 days

Warnings

Allergy alert: Do not use if you are allergic to polyethylene glycol

Do not use

if you have kidney disease, except under the advice and supervision of a doctor

Ask a doctor before use if you have

- nausea, vomiting or abdominal pain
- a sudden change in bowel habits that lasts over 2 weeks
- irritable bowel syndrome

Ask a doctor or pharmacist before use if you are

taking a prescription drug

When using this product

you may have loose, watery, more frequent stools

Stop use and ask a doctor if

- you have rectal bleeding or your nausea, bloating, cramping or abdominal pain gets worse. These may be signs of a serious condition.
- you get diarrhea
- you need to use a laxative for longer 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

- **do not take more than directed unless advised by your doctor**
- the bottle top is a measuring cap marked to contain 17 grams of powder when filled to the indicated line (white section in cap)
- adults and children 17 years of age and older:
- fill to top of white section in cap which is marked to indicate the correct dose (17 g)
- stir and dissolve in any 4 to 8 ounces of beverage (cold, hot or room temperature) then drink
- ensure that the powder is fully dissolved before drinking
- do not drink if there are any clumps
- use once a day
- use no more than 7 days
- children 16 years of age or under: ask a doctor

Other information

- store at 20°-25°C (68°-77°F)
- tamper-evident: do not use if printed foil seal under cap is missing, open or broken

Inactive ingredients

none

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to MiraLAX® active ingredient

Clearlax®

Polyethylene Glycol 3350 Powder for Solution

Osmotic Laxative

Relieves Occasional Constipation/Irregularity

Softens Stool

Unflavored Powder

Grit Free

NET WT. 8.3 OZ (235g)

14 ONCE-DAILY DOSES



Peel Here

Polyethylene Glycol 3350 increases frequency of bowel movements and softens the stool.
Dissolves in Any Beverage

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GLUTEN FREE

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Drug Facts (continued)

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)

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Inactive ingredients none

Questions or comments? 1-800-719-9260

• Sugar Free

*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark MiraLAX®.

MADE WITH PRIDE & CARE FOR H-E-B®
SAN ANTONIO, TX 78204

100% GUARANTEE promise

CLEAR LAX

polyethylene glycol 3350 powder, for solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-306
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P) (POLYETHYLENE GLYCOL 3350 - UNII:G2M7P15E5P)	POLYETHYLENE GLYCOL 3350	17 g in 17 g
Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			
Packaging			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-306-03	510 g in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2009	
2	NDC:37808-306-01	119 g in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2009	
3	NDC:37808-306-02	238 g in 1 BOTTLE; Type 0: Not a Combination Product	10/08/2009	
4	NDC:37808-306-18	153 g in 1 BOTTLE; Type 0: Not a Combination Product	08/16/2013	01/29/2020
5	NDC:37808-306-17	289 g in 1 BOTTLE; Type 0: Not a Combination Product	08/16/2013	
6	NDC:37808-306-19	595 g in 1 BOTTLE; Type 0: Not a Combination Product	07/31/2013	05/20/2021
7	NDC:37808-306-04	850 g in 1 BOTTLE; Type 0: Not a Combination Product	09/04/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090685	10/08/2009	

Labeler - H E B (007924756)

Revised: 3/2021

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