

**GAVILAX- polyethylene glycol 3350 powder, for solution**  
**Preferred Pharmaceuticals, Inc.**

-----  
**Drug Facts**

**Active Ingredient**

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 the 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.
- adults and children 17 years of age and older:
  - fill to top of line 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
- 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 Ingredient**

none

## **Questions or comments?**

**1-866-403-7592**

**Relabeled by Preferred Pharmaceuticals, Inc.**

## **PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

**NDC 68788-7508-02**

Original Prescription Strength

### **GaviLax**

Polyethylene Glycol 3350

Powder for Solution, Osmotic Laxative

**14 ONCE-DAILY DOSES**

**NET WT 8.3 OZ (238 g)**

# GaviLAX

Generic for Miralax

Each gm contains Polyethylene Glycol 3350

**Pkg Size:** Exp Date: #####  
Lot#: Batch#:

Ins:  
Mfg: Lupin Pharmaceuticals, Inc.  
Prod#:

### Warning

Store at 20°-25°C (68°-77°F). See bottle for directions. In case of overdose, get medical help or contact a Poison Control Center right away. If pregnant or breast-feeding, ask a health professional before use. Do not use if printed foil seal under cap is missing, open or broken. Do not use if you are allergic to polyethylene glycol. Keep this and all medication out of the reach of children.



CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.

GaviLAX  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Log



Directions English

Use as directed on package.  
Use as directed by your doctor



GTIN

#####  
SN #####  
EXP #####

Instrucciones Espanol:  
Utilice como dirigido en el paquete.  
Uso según lo dirigido por su doctor

GaviLAX  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Chart

GaviLAX  
Qty:  
Insurance NDC:  
Lot: Bat:

Billing

GaviLAX  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Patient

## GAVILAX

polyethylene glycol 3350 powder, for solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-7508(NDC:43386-312)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>POLYETHYLENE GLYCOL 3350</b> (UNII: G2M7P15E5P) (POLYETHYLENE GLYCOL 3350 - UNII:G2M7P15E5P)	POLYETHYLENE GLYCOL 3350	17 g in 17 g

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	

### Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7508-2	238 g in 1 BOTTLE; Type 0: Not a Combination Product	10/28/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA091077	10/09/2009	

**Labeler** - Preferred Pharmaceuticals, Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals, Inc. (791119022)

## Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals, Inc.		791119022	RELABEL(68788-7508)

Revised: 5/2026

Preferred Pharmaceuticals, Inc.