# GAVILAX - polyethylene glycol 3350 powder, for solution Lupin Pharmaceuticals, Inc.

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# **Drug Facts**

# **Active Ingredient**

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

# **Purpose**

#### 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:
- use once a day
- 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
- do not combine with starch-based thickeners used for difficulty swallowing
- ensure that the powder is fully dissolved before drinking
- do not drink if there are any clumps
- do not use 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, printed with "SEALED for YOUR PROTECTION" is missing, open or broken

# **Inactive Ingredient**

none

### Questions or comments?

#### 1-866-403-7592

- Dissolves in any beverages
- Sugar Free

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

**NDC** 43386-312-08

Original Prescription Strength

#### **GaviLax**

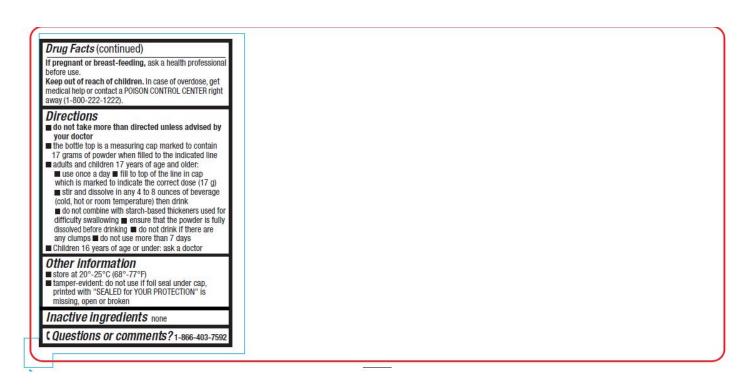
Polyethylene Glycol 3350

Powder for Oral Solution, Osmotic Laxative

14 ONCE-DAILY DOSES

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





**NDC** 43386-312-85

Original Prescription Strength

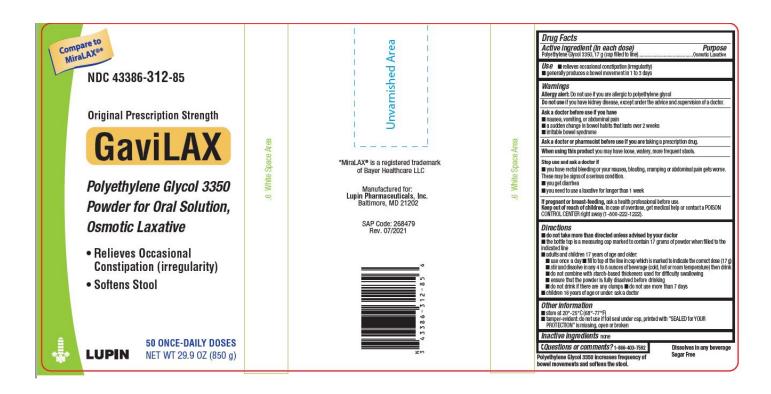
#### **GaviLax**

Polyethylene Glycol 3350

Powder for Oral Solution, Osmotic Laxative

**50 ONCE-DAILY DOSES** 

NET WT 29.9 OZ (850 g)



**NDC** 43386-312-07

Original Prescription Strength

#### **GaviLax**

Polyethylene Glycol 3350

Powder for Oral Solution, Osmotic Laxative
7 ONCE-DAILY DOSES

NET WT 4.1 OZ (119 g)



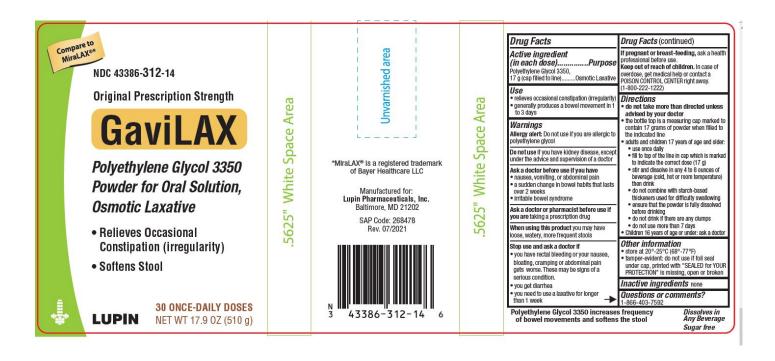
**NDC** 43386-312-14

Original Prescription Strength

#### GaviLax

Polyethylene Glycol 3350

# Powder for Oral Solution, Osmotic Laxative 30 ONCE-DAILY DOSES NET WT 17.9 OZ (510 g)



# **GAVILAX**

polyethylene glycol 3350 powder, for solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43386-312	
Route of Administration	ORAL			

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P) (POLYETHYLENE GLYCOL 3350 - UNII:G2M7P15E5P) POLYETHYLENE GLYCOL 3350 In 17 g in 17 g

Inactive Ingredients	
Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Product Characteristics		
Color	WHITE	Score
Shape		Size
Flavor		Imprint Code

#### **Contains**

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43386-312- 14	510 g in 1 BOTTLE; Type 0: Not a Combination Product	10/09/2009	
2	NDC:43386-312- 08	238 g in 1 BOTTLE; Type 0: Not a Combination Product	10/09/2009	
3	NDC:43386-312- 07	119 g in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2040	
4	NDC:43386-312- 85	850 g in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2040	

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

# Labeler - Lupin Pharmaceuticals,Inc. (089153071)

# **Registrant -** Novel Laboratories, Inc. (793518643)

Establishment				
Name	Address	ID/FEI	Business Operations	
Novel Laboratories, Inc.		793518643	ANALYSIS (43386-312), MANUFACTURE (43386-312), PACK (43386-312)	

Revised: 1/2024 Lupin Pharmaceuticals,Inc.