UP AND UP POWDERLAX- polyethylene glycol 3350 powder, for solution
Target Corporation

Target Corporation Powderlax® 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:
- use once a day
- 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
- 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 is missing, open or broken

Inactive ingredients

none

Questions or comments?

1-888-547-7400

Principal Display Panel

Compare to active ingredient in MiraLAX®

Powderlax®

polyethylene glycol 3350 powder for solution, osmotic laxative

relieves occasional constipation/irregularity

softens stool

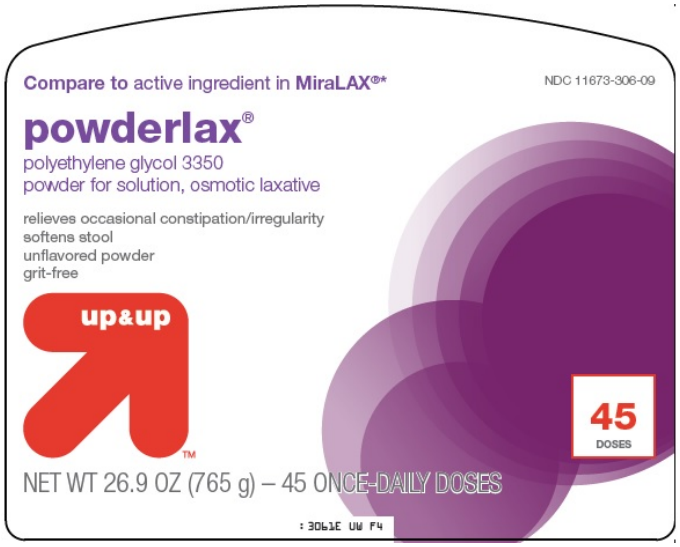
unflavored powder

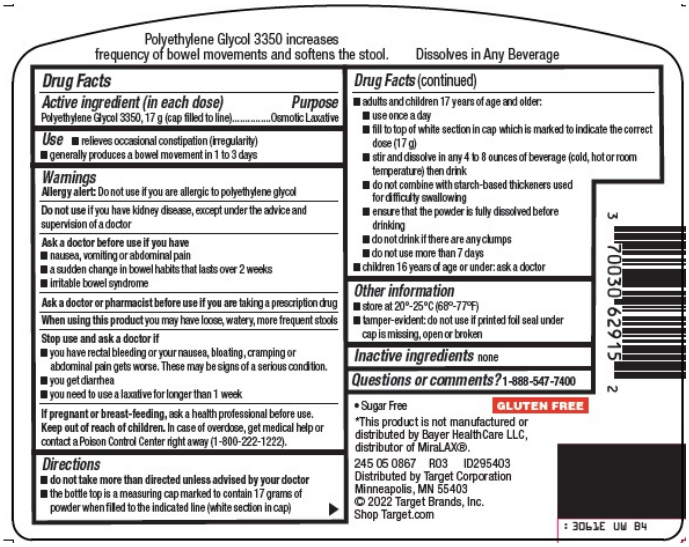
grit-free

up&up™

45 DOSES

NET WT 26.9 OZ (765 g) – 45 ONCE-DAILY DOSES





UP AND UP POWDERLAX

polyethylene glycol 3350 powder, for solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-306
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				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-306-02	238 g in 1 BOTTLE; Type 0: Not a Combination Product	10/07/2009	

2	NDC:11673-306-03	510 g in 1 BOTTLE; Type 0: Not a Combination Product	10/07/2009	
3	NDC:11673-306-19	595 g in 1 BOTTLE; Type 0: Not a Combination Product	10/09/2013	02/03/2016
4	NDC:11673-306-04	850 g in 1 BOTTLE; Type 0: Not a Combination Product	03/03/2014	07/23/2019
5	NDC:11673-306-17	289 g in 1 BOTTLE; Type 0: Not a Combination Product	08/27/2014	10/28/2016
6	NDC:11673-306-09	765 g in 1 BOTTLE; Type 0: Not a Combination Product	10/21/2016	
7	NDC:11673-306-01	119 g in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2019	
8	NDC:11673-306-52	10 in 1 CARTON	04/11/2019	11/01/2020
8		17 g in 1 PACKET; Type 0: Not a Combination Product		
9	NDC:11673-306-60	20 in 1 CARTON	04/11/2019	12/01/2020
9		17 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090685	10/07/2009	

Labeler - Target Corporation (006961700)

Revised: 2/2023 Target Corporation