

EZ2GO - polyethylene glycol 3350 powder
Valley Medical Products,LLC

Inactive Ingredients none

Active Ingredient

Polyethylene Glycol 3350, 17 g

Purpose

Osmotic Laxative

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

When using this product, you may have loose, watery, more frequent stools

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

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
- side effects may occur. you may report the side effects

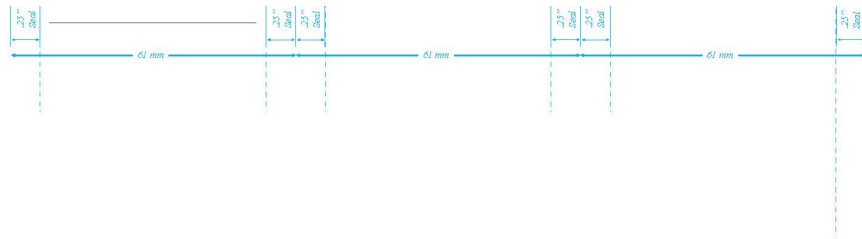
to FDA at 1 - 800 - FDA - 1088

Keep out of reach of children. 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.

Directions

- do not take more than directed unless advised by your doctor
- adults and children 17 years of age and older:
 - stir and dissolve one packet (17g) in any 4 to 8 ounces of beverage (cold, hot or room temperature) and then drink
 - use once a day
 - use no more than 7 days
- Children 16 years of age or under: ask a doctor



EZ2GO

polyethylene glycol 3350 powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76470-005
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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76470-005-05	238 g in 1 BOTTLE; Type 0: Not a Combination Product	10/20/2012	
2	NDC:76470-005-17	17 g in 1 POUCH; Type 0: Not a Combination Product	10/20/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA091077	10/20/2012	

Labeler - Valley Medical Products,LLC (969389407)

Establishment

Name	Address	ID/FEI	Business Operations
Valley Medical Products,LLC		969389407	relabel(76470-005) , repack(76470-005)

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Revised: 1/2017

Valley Medical Products,LLC