

Drugs@FDA

FDA Approved Drug Products

[Start Over](#)[Back to Overview](#)[FAQ](#) | [Instructions](#) | [Glossary](#) | [Contact Us](#)**Drug Details**

Drug Name(s)	GLYCOLAX (Generic Drug)
FDA Application No.	(ANDA) 090600
Active Ingredient(s)	POLYETHYLENE GLYCOL 3350
Company	KREMERS URBAN PHARMS
Original Approval or Tentative Approval Date	October 6, 2009

- [Other OTC Drugs with the same Active Ingredient, Strength and Dosage Form/Route](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- [Label Information](#)

Products on Application (ANDA) #090600

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form / Route	Marketing Status	RLD	TE Code
GLYCOLAX	POLYETHYLENE GLYCOL 3350	17GM/PACKET	FOR SOLUTION; ORAL	Over-the-counter	No	None
GLYCOLAX	POLYETHYLENE GLYCOL 3350	17GM/SCOOPFUL	FOR SOLUTION; ORAL	Over-the-counter	No	None

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)[Disclaimer](#)

FDA/Center for Drug Evaluation and Research
Office of Training and Communications
Division of Information Services
Update Frequency: Daily