Visited 03/8/2011

Drug Details

Drug Name(s)	**NORCO (Generic Drug)**
FDA Application No.	**(ANDA) 040099**
Active Ingredient(s)	**ACETAMINOPHEN; HYDROCODONE BITARTRATE**
Company	**WATSON LABS**
Original Approval or Tentative Approval Date	**June 25, 1997**

- Therapeutic Equivalents
- Approval History, Letters, Reviews, and Related Documents
- **Labels are not available**

Products on Application (ANDA) #040099
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NORCO	ACETAMINOPHEN; HYDROCODONE BITARTRATE	325MG; 5MG	TABLET; ORAL	Prescription	Yes	AA

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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