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Drug Details

Drug Name(s)	CARBINOXAMINE MALEATE
FDA Application No.	(ANDA) 207484
Active Ingredient(s)	CARBINOXAMINE MALEATE
Company	MIKART INC
Original Approval or Tentative Approval Date	May 31, 2016

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (ANDA) #207484

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
CARBINOXAMINE MALEATE	CARBINOXAMINE MALEATE	6MG	TABLET;ORAL	Prescription	No	None

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