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Drug Details

Drug Name(s)	CHLORZOAZONE
FDA Application No.	(ANDA) 207483
Active Ingredient(s)	CHLORZOAZONE
Company	MIKART INC
Original Approval or Tentative Approval Date	June 24, 2016

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

Products on Application (ANDA) #207483

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
CHLORZOAZONE	CHLORZOAZONE	250MG	TABLET;ORAL	Prescription	Yes	None

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