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Drug Details

Drug Name(s)	OXYBUTYNIN CHLORIDE
FDA Application No.	(ANDA) 207138
Active Ingredient(s)	OXYBUTYNIN CHLORIDE
Company	ACCORD HLTHCARE
Original Approval or Tentative Approval Date	February 29, 2016

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #207138

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
OXYBUTYNIN CHLORIDE	OXYBUTYNIN CHLORIDE	5MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYBUTYNIN CHLORIDE	OXYBUTYNIN CHLORIDE	10MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
OXYBUTYNIN CHLORIDE	OXYBUTYNIN CHLORIDE	15MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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U.S. Department of **Health & Human Services**

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