Drug Details

Drug Name(s)	**ULTRAVATE**
FDA Application No.	**(NDA) 208183**
Active Ingredient(s)	**HALOBETASOL PROPIONATE**
Company	**FERNDALE LABS**
Original Approval or Tentative Approval Date	**November 6, 2015**
Chemical Type	**3 New dosage form**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (NDA) #208183
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ULTRAVATE	HALOBETASOL PROPIONATE	0.05%	LOTION;TOPICAL	Prescription	TBD [11]	None

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