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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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