Drug Details

Drug Name(s)	**DEXMETHYLPHENIDATE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 204534**
Active Ingredient(s)	**DEXMETHYLPHENIDATE HYDROCHLORIDE**
Company	**NOVEL LABS INC**

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

Products on Application (ANDA) #204534
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DEXMETHYLPHENIDATE HYDROCHLORIDE	DEXMETHYLPHENIDATE HYDROCHLORIDE	2.5MG	TABLET;ORAL	None (Tentative Approval)	No	None
DEXMETHYLPHENIDATE HYDROCHLORIDE	DEXMETHYLPHENIDATE HYDROCHLORIDE	5MG	TABLET;ORAL	None (Tentative Approval)	No	None
DEXMETHYLPHENIDATE HYDROCHLORIDE	DEXMETHYLPHENIDATE HYDROCHLORIDE	10MG	TABLET;ORAL	None (Tentative Approval)	No	None

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