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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	RLDTE 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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