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Drug Details

Drug Name(s)	METHYLPHENIDATE HYDROCHLORIDE
FDA Application No.	(ANDA) 076655
Active Ingredient(s)	METHYLPHENIDATE HYDROCHLORIDE
Company	ACTAVIS LABS FL INC

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (ANDA) #076655
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	18MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	27MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	36MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	54MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None

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