Drug Details

Drug Name(s)	**METHYLPHENIDATE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 206726**
Active Ingredient(s)	**METHYLPHENIDATE HYDROCHLORIDE**
Company	**MYLAN PHARMS INC**
Original Approval or Tentative Approval Date	**October 21, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #206726
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	18MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	27MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	36MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	54MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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