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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

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- **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	RLDTE 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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