Drug Details

Drug Name(s)	**METHYLPHENIDATE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 203583**
Active Ingredient(s)	**METHYLPHENIDATE HYDROCHLORIDE**
Company	**MALLINCKRODT INC**
Original Approval or Tentative Approval Date	**September 29, 2015**

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

Products on Application (ANDA) #203583
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	10MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	20MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	30MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	40MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	50MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2
METHYLPHENIDATE HYDROCHLORIDE	METHYLPHENIDATE HYDROCHLORIDE	60MG	CAPSULE, EXTENDED RELEASE;ORAL	Prescription	No	AB2

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