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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	RLDTE 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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