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Drug Details

Drug Name(s)	METHOCARBAMOL
FDA Application No.	(ANDA) 205354
Active Ingredient(s)	METHOCARBAMOL
Company	MONTEREY PHARMS LLC
Original Approval or Tentative Approval Date	October 27, 2016

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Products on Application (ANDA) #205354

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
METHOCARBAMOL	METHOCARBAMOL	1GM/10ML (100MG/ML)	SOLUTION;IM-IV	Prescription	No AP

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