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Drug Details

Drug Name(s)	BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE
FDA Application No.	(ANDA) 201633
Active Ingredient(s)	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE
Company	SUN PHARM INDS LTD
Original Approval or Tentative Approval Date	August 5, 2016

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- **Labels are not available**

Products on Application (ANDA) #201633

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 2MG BASE; EQ 0.5MG BASE	TABLET;SUBLINGUAL	Prescription No	AB
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 8MG BASE; EQ 2MG BASE	TABLET;SUBLINGUAL	Prescription No	AB

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