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**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **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	RLD	TE 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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