Drug Details

Drug Name(s)	**BUNAVAIL**
FDA Application No.	**(NDA) 205637**
Active Ingredient(s)	**BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE**
Company	**BIODELIVERY SCI INTL**
Original Approval or Tentative Approval Date	**June 6, 2014**
Chemical Type	**3 New dosage form**
Review Classification	**S Standard review drug**

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**
- **REMS[11]**

Products on Application (NDA) #205637
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BUNAVAIL	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 2.1MG BASE; EQ 0.3MG BASE	FILM;BUCCAL	Prescription	No	None
BUNAVAIL	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 4.2MG BASE; EQ 0.7MG BASE	FILM;BUCCAL	Prescription	No	None
BUNAVAIL	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 6.3MG BASE; EQ 1MG BASE	FILM;BUCCAL	Prescription	Yes	None

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