Drug Details

Drug Name(s)	**SUBOXONE**
FDA Application No.	**(NDA) 020733**
Active Ingredient(s)	**BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE**
Company	**RECKITT BENCKISER**
Original Approval or Tentative Approval Date	**October 8, 2002**
Chemical Type	**4 New combination**
Review Classification	**P Priority review drug** **O Orphan drug**

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Other Important Information from FDA** [11]
- **Label Information**
- **REMS**[10]
- **Medication Guide**[12]

Products on Application (NDA) #020733
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
SUBOXONE	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 2MG BASE; EQ 0.5MG BASE	TABLET;SUBLINGUAL	Discontinued	No	None
SUBOXONE	BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE	EQ 8MG BASE; EQ 2MG BASE	TABLET;SUBLINGUAL	Discontinued	No	None

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