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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](#)
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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	RLDTE 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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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)
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