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Drug Details

Drug Name(s)	BUTRANS
FDA Application No.	(NDA) 021306
Active Ingredient(s)	BUPRENORPHINE
Company	PURDUE PHARMA LP
Original Approval or Tentative Approval Date	June 30, 2010
Chemical Type	3 New dosage form
Review Classification	S Standard review drug

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #021306

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BUTRANS	BUPRENORPHINE	5MCG/HR	FILM, EXTENDED RELEASE;TRANSDERMAL	Prescription	No	None
BUTRANS	BUPRENORPHINE	10MCG/HR	FILM, EXTENDED RELEASE;TRANSDERMAL	Prescription	No	None
BUTRANS	BUPRENORPHINE	20MCG/HR	FILM, EXTENDED RELEASE;TRANSDERMAL	Prescription	Yes	None
BUTRANS	BUPRENORPHINE	15MCG/HR	FILM, EXTENDED RELEASE;TRANSDERMAL	Prescription	No	None
BUTRANS	BUPRENORPHINE	7.5MCG/HR	FILM, EXTENDED RELEASE;TRANSDERMAL	Prescription	No	None

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