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## Drug Details

<b>Drug Name(s)</b>	<b>HYSINGLA</b>
<b>FDA Application No.</b>	<b>(NDA) 206627</b>
<b>Active Ingredient(s)</b>	<b>HYDROCODONE BITARTRATE</b>
<b>Company</b>	<b>PURDUE PHARMA LP</b>
<b>Original Approval or Tentative Approval Date</b>	<b>November 20, 2014</b>
<b>Chemical Type</b>	<b>3 New dosage form</b>
<b>Review Classification</b>	<b>P Priority review drug</b>

- [There are no Therapeutic Equivalents](#)
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### Products on Application (NDA) #206627

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HYSINGLA	HYDROCODONE BITARTRATE	20MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	Yes	None
HYSINGLA	HYDROCODONE BITARTRATE	30MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None
HYSINGLA	HYDROCODONE BITARTRATE	40MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None
HYSINGLA	HYDROCODONE BITARTRATE	60MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None
HYSINGLA	HYDROCODONE BITARTRATE	80MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None
HYSINGLA	HYDROCODONE BITARTRATE	100MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None
HYSINGLA	HYDROCODONE BITARTRATE	120MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	None

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