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Drug Details

Drug Name(s)	ENTECAVIR
FDA Application No.	(ANDA) 206652
Active Ingredient(s)	ENTECAVIR
Company	AMNEAL PHARMS
Original Approval or Tentative Approval Date	November 12, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- Labels are not available

Products on Application (ANDA) #206652

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ENTECAVIR	ENTECAVIR	0.5MG	TABLET;ORAL	Prescription	No	AB
ENTECAVIR	ENTECAVIR	1MG	TABLET;ORAL	Prescription	No	AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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