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Drug Details

Drug Name(s)	ABACAVIR SULFATE
FDA Application No.	(ANDA) 201107
Active Ingredient(s)	ABACAVIR SULFATE
Company	HETERO LABS LTD III
Original Approval or Tentative Approval Date	September 26, 2016

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

Products on Application (ANDA) #201107

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ABACAVIR SULFATE	ABACAVIR SULFATE	EQ 20MG BASE/ML	SOLUTION;ORAL	Prescription	No	AA

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

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