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Drug Details

Drug Name(s)	DEXTROSE 5% IN PLASTIC CONTAINER
FDA Application No.	(ANDA) 207449
Active Ingredient(s)	DEXTROSE
Company	FRESENIUS KABI USA
Original Approval or Tentative Approval Date	October 21, 2016

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- **Labels are not available**

Products on Application (ANDA) #207449
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DEXTROSE 5% IN PLASTIC CONTAINER	DEXTROSE	50MG/ML	INJECTABLE;INJECTION	Prescription	No AP

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