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Drug Details

Drug Name(s)	SODIUM POLYSTYRENE SULFONATE
FDA Application No.	(ANDA) 206815
Active Ingredient(s)	SODIUM POLYSTYRENE SULFONATE
Company	INVATECH PHARMA
Original Approval or Tentative Approval Date	February 18, 2016

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

Products on Application (ANDA) #206815

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
SODIUM POLYSTYRENE SULFONATE	SODIUM POLYSTYRENE SULFONATE	454GM/BOT	POWDER;ORAL, RECTAL	Prescription No	AA

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