Drug Details

Drug Name(s)	**SODIUM POLYSTYRENE SULFONATE**
FDA Application No.	**(ANDA) 205727**
Active Ingredient(s)	**SODIUM POLYSTYRENE SULFONATE**
Company	**BELCHER PHARMS LLC**
Original Approval or Tentative Approval Date	**February 23, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #205727
Click on a column header to re-sort the table:

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

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