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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

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- **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	RLDTE Code
SODIUM POLYSTYRENE SULFONATE	SODIUM POLYSTYRENE SULFONATE	454GM/BOT	POWDER;ORAL, RECTAL	Prescription No	AA

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U.S. Food and Drug Administration
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